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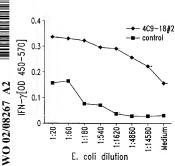
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(54) Title: COMPOUNDS AND METHODS FOR TREATMENT AND DIAGNOSIS OF CHLAMYDIAL INFECTION



(57) Abstract: Compounds and methods for the diagnosis and treatment of Chlamydial infection are disclosed. The compounds provided include polypeptides that contain at least one antigenic portion of a Chlamvdia antigen and DNA sequences encoding such polypeptides. Pharmaceutical compositions and vaccines comprising such polypeptides or DNA sequences are also provided, together with antibodies directed against such polypeptides. Diagnostic kits containing such polypeptides or DNA sequences and a suitable detection reagent may be used for the detection of Chlamydial infection in patients and in biological samples.

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COMPOUNDS AND METHODS FOR TREATMENT AND DIAGNOSIS OF CHLAMYDIAL INFECTION

TECHNICAL FIELD

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The present invention relates generally to the detection and treatment of Chlamydial infection. In particular, the invention is related to polypeptides comprising a *Chlamydia* antigen and the use of such polypeptides for the serodiagnosis and treatment of Chlamydial infection.

10 BACKGROUND OF THE INVENTION

Chlamydiae are intracellular bacterial pathogens that are responsible for a wide variety of important human and animal infections. Chlamydia trachomatis is one of the most common causes of sexually transmitted diseases and can lead to pelvic inflammatory disease (PID), resulting in tubal obstruction and infertility. Chlamydia trachomatis may also play a role in male infertility. In 1990, the cost of treating PID in the US was estimated to be \$4 billion. Trachoma, due to ocular infection with Chlamydia trachomatis, is the leading cause of preventable blindness worldwide. Chlamydia pneumonia is a major cause of acute respiratory tract infections in humans and is also believed to play a role in the pathogenesis of atherosclerosis and, in particular, coronary heart disease. Individuals with a high titer of antibodies to Chlamydia pneumonia have been shown to be at least twice as likely to suffer from coronary heart disease as seronegative individuals. Chlamydial infections thus constitute a significant health problem both in the US and worldwide.

Chlamydial infection is often asymptomatic. For example, by the time a woman seeks medical attention for PID, irreversible damage may have already occurred resulting in infertility. There thus remains a need in the art for improved vaccines and pharmaceutical compositions for the prevention and treatment of *Chlamydia* infections. The present invention fulfills this need and further provides other related advantages.

0 SUMMARY OF THE INVENTION

The present invention provides compositions and methods for the diagnosis and therapy of *Chlamydia* infection. In one aspect, the present invention

provides polypeptides comprising an immunogenic portion of a *Chlamydia* antigen, or a variant of such an antigen. Certain portions and other variants are immunogenic, such that the ability of the variant to react with antigen-specific antisera is not substantially diminished. Within certain embodiments,, the polypeptide comprises an amino acid sequence encoded by a polynucleotide sequence selected from the group consisting of (a) a sequence of SEQ ID NO: 358-361, 366-385, 406-430, 455-489, 516-517, 523-559, and 582-596; (b) the complements of said sequences; and (c) sequences that hybridize to a sequence of (a) or (b) under moderate to highly stringent conditions. In specific embodiments, the polypeptides of the present invention comprise at least a portion of a *Chlamydial* protein that includes an amino acid sequence selected from the group consisting of sequences recited in SEQ ID NO:362-365, 386-405, 431-454, 490-515, 518-522, 560-581, and 597-599 and variants thereof.

The present invention further provides polynucleotides that encode a
polypeptide as described above, or a portion thereof (such as a portion encoding at least
15 amino acid residues of a *Chlamydial* protein), expression vectors comprising such
polynucleotides and host cells transformed or transfected with such expression vectors.

In a related aspect, polynucleotide sequences encoding the above polypeptides, recombinant expression vectors comprising one or more of these polynucleotide sequences and host cells transformed or transfected with such expression vectors are also provided.

In another aspect, the present invention provides fusion proteins comprising an inventive polypeptide, or, alternatively, an inventive polypeptide and a known *Chlamydia* antigen, as well as polynucleotides encoding such fusion proteins, in combination with a physiologically acceptable carrier or immunostimulant for use as pharmaceutical compositions and vaccines thereof.

The present invention further provides pharmaceutical compositions that comprise: (a) an antibody, both polyclonal and monoclonal, or antigen-binding fragment thereof that specifically binds to a *Chlamydial* protein; and (b) a physiologically acceptable carrier. Within other aspects, the present invention provides opharmaceutical compositions that comprise one or more *Chlamydia* polypeptides disclosed herein, e.g., a polypeptide according to SEQ ID NO:362-365, 386-405, 431-454, 490-515, 518-522, 560-581, and 597-599, or a polynucleotide molecule encoding

such a polypeptide, such as a polynucleotide according to SEO ID NO:358-361, 366-385, 406-430, 455-489, 516-517, 523-559, and 582-596, and a physiologically acceptable carrier. The invention also provides vaccines for prophylactic and therapeutic purposes comprising one or more of the disclosed polypeptides and an immunostimulant, as defined herein, together with vaccines comprising one or more polynucleotide sequences encoding such polypeptides and an immunostimulant.

In yet another aspect, methods are provided for inducing protective immunity in a patient, comprising administering to a patient an effective amount of one or more of the above pharmaceutical compositions or vaccines.

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In yet a further aspect, methods for the treatment of Chlamydia infection in a patient are provided, the methods comprising obtaining peripheral blood mononuclear cells (PBMC) from the patient, incubating the PBMC with a polypeptide of the present invention (or a polynucleotide that encodes such a polyneptide) to provide incubated T cells and administering the incubated T cells to the patient. The present 15 invention additionally provides methods for the treatment of Chlamvdia infection that comprise incubating antigen presenting cells with a polypeptide of the present invention (or a polynucleotide that encodes such a polypeptide) to provide incubated antigen presenting cells and administering the incubated antigen presenting cells to the patient. Proliferated cells may, but need not, be cloned prior to administration to the patient. In certain embodiments, the antigen presenting cells are selected from the group consisting of dendritic cells, macrophages, monocytes, B-cells, and fibroblasts. Compositions for the treatment of Chlamvdia infection comprising T cells or antigen presenting cells that have been incubated with a polypeptide or polynucleotide of the present invention are also provided. Within related aspects, vaccines are provided that comprise: (a) an antigen presenting cell that expresses a polypeptide as described above and (b) an immunostimulant.

The present invention further provides, within other aspects, methods for removing Chlamydial-infected cells from a biological sample, comprising contacting a biological sample with T cells that specifically react with a Chlamydial protein, wherein the step of contacting is performed under conditions and for a time sufficient to permit the removal of cells expressing the protein from the sample.

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Within related aspects, methods are provided for inhibiting the development of Chlamydial infection in a patient, comprising administering to a patient a biological sample treated as described above. In further aspects of the subject invention, methods and diagnostic kits are provided for detecting Chlamydia infection in a patient. In one embodiment, the method comprises: (a) contacting a biological sample with at least one of the polypeptides or fusion proteins disclosed herein; and (b) detecting in the sample the presence of binding agents that bind to the polypeptide or fusion protein, thereby detecting Chlamydia infection in the biological sample. Suitable biological samples include whole blood, sputum, serum, plasma, saliva, cerebrospinal 10 fluid and urine. In one embodiment, the diagnostic kits comprise one or more of the polypeptides or fusion proteins disclosed herein in combination with a detection reagent. In yet another embodiment, the diagnostic kits comprise either a monoclonal antibody or a polyclonal antibody that binds with a polypeptide of the present invention.

The present invention also provides methods for detecting Chlamydia 15 infection comprising: (a) obtaining a biological sample from a patient; (b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, at least one of the oligonucleotide primers being specific for a polynucleotide sequence disclosed herein; and (c) detecting in the sample a polynucleotide sequence that amplifies in the presence of the oligonucleotide primers. In one embodiment, the oligonucleotide primer comprises at least about 10 contiguous nucleotides of a polynucleotide sequence peptide disclosed herein, or of a sequence that hybridizes thereto.

In a further aspect, the present invention provides a method for detecting Chlamydia infection in a patient comprising: (a) obtaining a biological sample from the 25 patient; (b) contacting the sample with an oligonucleotide probe specific for a polynucleotide sequence disclosed herein; and (c) detecting in the sample a polynucleotide sequence that hybridizes to the oligonucleotide probe. In one embodiment, the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide sequence disclosed herein, or a sequence that hybridizes thereto.

These and other aspects of the present invention will become apparent upon reference to the following detailed description. All references disclosed herein are

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hereby incorporated by reference in their entirety as if each was incorporated individually.

SEQUENCE IDENTIFIERS

SEQ ID NO: 1 is the determined DNA sequence for the *C. trachomatis* clone 1-R1-66

SEQ ID NO: 2 is the determined DNA sequence for the *C. trachomatis* clone 4-D7-28.

SEQ ID NO: 3 is the determined DNA sequence for the C. trachomatis

clone 3-G3-10.

SEQ ID NO: 4 is the determined DNA sequence for the *C. trachomatis* clone 10-C10-31.

SEQ ID NO: 5 is the predicted amino acid sequence for 1-B1-66.

SEQ ID NO: 6 is the predicted amino acid sequence for 4-D7-28.

SEQ ID NO: 7 is a first predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 8 is a second predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 9 is a third predicted amino acid sequence for 3-G3-10.

SEO ID NO: 10 is a fourth predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 11 is a fifth predicted amino acid sequence for 3-G3-10.

SEQ ID NO: 12 is the predicted amino acid sequence for 10-C10-31.

SEQ ID NO: 13 is the amino acid sequence of the synthetic peptide 1-B1-66/48-67

SEQ ID NO: 14 is the amino acid sequence of the synthetic peptide 1-B1-66/58-77.

SEQ ID NO: 15 is the determined DNA sequence for the *C. trachomatis* serovar LGV II clone 2C7-8

SEQ ID NO: 16 is a DNA sequence of a putative open reading frame from a region of the C. trachomatis serovar D genome to which 2C7-8 maps

SEQ ID NO: 17 is the predicted amino acid sequence encoded by the

DNA sequence of SEQ ID NO: 16

SEQ ID NO: 18 is the amino acid sequence of the synthetic peptide
CrC7.8-12

SEQ ID NO: 19 is the amino acid sequence of the synthetic peptide CtC7.8-13

SEQ ID NO: 20 is the predicted amino acid sequence encoded by a second putative open reading from C. trachomatis serovar D

5 SEQ ID NO: 21 is the determined DNA sequence for clone 4C9-18 from C. trachomatis LGV II

SEQ ID NO: 22 is the determined DNA sequence homologous to Lipoamide Dehvdrogenase from C. trachomatis LGV II

SEQ ID NO: 23 is the determined DNA sequence homologous to

Hypothetical protein from C. trachomatis LGV II

SEQ ID NO: 24 is the determined DNA sequence homologous to Ubiquinone Mehtyltransferase from C. trachomatis LGV II

SEQ ID NO: 25 is the determined DNA sequence for clone 4C9-18#2 BL21 pLysS from C. trachomatis LGV II

15 SEQ ID NO: 26 is the predicted amino acid sequence for 4C9-18#2 from C. trachomatis LGV II

SEQ ID NO: 27 is the determined DNA sequence for Cp-SWIB from C. pneumonia strain TWAR

SEQ ID NO: 28 is the predicted amino acid sequence for Cp-SWIB from C, pneumonia strain TWAR

SEQ ID NO: 29 is the determined DNA sequence for Cp-S13 (CT509) from C. pneumonia strain TWAR

SEQ ID NO: 30 is the predicted amino acid sequence for Cp-S13 from $\it C.~pneumonia$ strain TWAR

25 SEQ ID NO: 31 is the amino acid sequence for a 10mer consensus peptide from CtC7.8-12 and CtC7.8-13

SEQ ID NO: 32 is the predicted amino acid sequence for clone 2C7-8 from C. trachonatis LGV II

SEQ ID NO: 33 is the DNA sequence corresponding to nucleotides
597304-597145 of the *C. trachomatis* serovar D genome (NCBI, BLASTN search),
which shows homology to clone 2C7-8

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SEQ ID NO: 34 is the predicted amino acid sequence encoded by the sequence of SEO ID NO: 33

SEQ ID NO: 35 is the DNA sequence for C.p. SWIB Nde (5' primer) from C. pneumonia

5 SEQ ID NO: 36 is the DNA sequence for C.p. SWIB EcoRI (3' primer) from C. pneumonia

SEQ ID NO : 37 is the DNA sequence for C.p. S13 Nde (5' primer) from $\emph{C. pneumonia}$

SEQ ID NO: 38 is the DNA sequence for C.p. S13 EcoR1 (3' primer) from C. pneumonia

SEQ ID NO: 39 is the amino acid sequence for CtSwib 52-67 peptide from C. trachomatis LGV II

SEQ ID NO: 40 is the amino acid sequence for CpSwib 53-68 peptide from C, pneumonia

SEQ ID NO: 41 is the amino acid sequence for HuSwib 288-302 peptide from Human SWI domain

SEQ ID NO: 42 is the amino acid sequence for CtSWI-T 822-837 peptide from the topoisomerase-SWIB fusion of C. trachomatis

SEQ ID NO: 43 is the amino acid sequence for CpSWI-T 828-842 peptide from the topoisomerase-SWIB fusion of *C. pneumonia*

SEQ ID NO: 44 is a first determined DNA sequence for the C. trachomatis LGV II clone 19783.3,jen.seq(1>509)CTL2#11-3', representing the 3' end.

SEQ ID NO: 45 is a second determined DNA sequence for the C. trachomatis LGV II clone 19783.4.ien.sea(1>481)CTL2#11-5', representing the 5' end.

SEQ ID NO: 46 is the determined DNA sequence for the *C. trachomatis* LGV II clone19784CTL2 12consensus.seq(1>427)CTL2#12.

SEQ ID NO: 47 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19785.4.jen.seq(1>600)CTL2#16-5', representing the 5' end.

SEQ ID NO: 48 is a first determined DNA sequence for the C. trachomatis LGV II clone 19786.3,jen.seq(1>600)CTL2#18-3', representing the 3' end.

SEQ ID NO: 49 is a second determined DNA sequence for the C. trachomatis LGV II clone 19786.4,jen.seq(1>600)CTL2#18-5', representing the 5' end.

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SEQ ID NO: 50 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19788CTL2 21consensus.sea(1>406)CTL2#21.

SEQ ID NO: 51 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19790CTL2 23consensus.seq(1>602)CTL2#23.

5 SEQ ID NO: 52 is the determined DNA sequence for the C. trachomatis LGV II clone 19791CTL2 24consensus.seq(1>145)CTL2#24.

SEQ ID NO: 53 is the determined DNA sequence for the *C. trachomatis* LGV II clone CTL2#4.

SEQ ID NO: 54 is the determined DNA sequence for the *C. trachomatis*10 LGV II clone CTL2#8b.

SEQ ID NO: 55 is the determined DNA sequence for the *C. trachomatis* LGV II clone15-G1-89, sharing homology to the lipoamide dehydrogenase gene CT557.

SEQ ID NO: 56 is the determined DNA sequence for the *C. trachomatis*LGV II clone 14-H1-4, sharing homology to the thiol specific antioxidant gene CT603.

SEQ ID NO: 57 is the determined DNA sequence for the *C. trachomatis* LGV II clone 12-G3-83, sharing homology to the hypothetical protein CT622.

SEQ ID NO: 58 is the determined DNA sequence for the *C. trachomatis* LGV II clone 12-B3-95, sharing homology to the lipoamide dehydrogenase gene CT557.

20 SEQ ID NO: 59 is the determined DNA sequence for the C. trachomatis LGV II clone 11-H4-28, sharing homology to the dnaK gene CT396.

SEQ ID NO: 60 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-H3-68, sharing partial homology to the PGP6-D virulence protein and L1 ribosomal gene CT318.

SEQ ID NO: 61 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-G1-34, sharing partial homology to the malate dehydrogenase gene CT376 and to the glycogen hydrolase gene CT042.

SEQ ID NO: 62 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-G10-46, sharing homology to the hypothetical protein CT610.

SEQ ID NO: 63 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-C12-91, sharing homology to the OMP2 gene CT443.

15 LGV II clone CtL2#2.

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SEQ ID NO: 64 is the determined DNA sequence for the *C. trachomatis* LGV II clone 11-A3-93, sharing homology to the HAD superfamily gene CT103.

SEQ ID NO: 65 is the determined amino acid sequence for the C. trachomatis LGV II clone 14-H1-4, sharing homology to the thiol specific antioxidant 5 gene CT603.

SEQ ID NO: 66 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#9.

SEQ ID NO: 67 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#7.

10 SEQ ID NO: 68 is the determined DNA sequence for the C. trachomatis LGV II clone Ctl 2#6.

SEQ ID NO: 69 is the determined DNA sequence for the *C. trachomatis* LGV II clone CtL2#5.

SEQ ID NO: 70 is the determined DNA sequence for the C. trachomatis

SEQ ID NO: 71 is the determined DNA sequence for the C. trachomatis
LGV II clone Ctl.2#1.

SEQ ID NO: 72 is a first determined DNA sequence for the C. trachomatis LGV II clone 23509.2CtL2#3-5', representing the 5' end.

20 SEQ ID NO: 73 is a second determined DNA sequence for the C. trachomatis LGV II clone 23509.1Ctl.2#3-3', representing the 3' end.

SEQ ID NO: 74 is a first determined DNA sequence for the *C. trachomatis* LGV II clone 22121.2Ctl.2#10-5', representing the 5' end.

SEQ ID NO: 75 is a second determined DNA sequence for the C. trachomatis LGV II clone 22121.1CiL2#10-3', representing the 3' end.

SEQ ID NO: 76 is the determined DNA sequence for the C. trachomatis

LGV II clone 19787.6Ctl.2#19-5', representing the 5' end.

SEO ID NO: 77 is the determined DNA sequence for the *C. pneumoniae*

SEQ ID NO: 77 is the determined DNA sequence for the *C. pneumoniae* LGV II clone CpS13-His.

30 SEQ ID NO: 78 is the determined DNA sequence for the C. pneumoniae LGV II clone Cp SWIB-His.

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SEQ ID NO: 79 is the determined DNA sequence for the *C. trachomatis* LGV II clone 23-G7-68, sharing partial homology to the L11, L10 and L1 ribosomal protein.

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SEQ ID NO: 80 is the determined DNA sequence for the *C. trachomatis*5 LGV II clone 22-F8-91, sharing homology to the pmpC gene.

SEQ ID NO: 81 is the determined DNA sequence for the *C. trachomatis* LGV II clone 21-E8-95, sharing homology to the CT610-CT613 genes.

SEQ ID NO: 82 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-F12-57, sharing homology to the CT858 and recA genes.

SEQ ID NO: 83 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-F12-53, sharing homology to the CT445 gene encoding glutamyl tRNA synthetase.

SEQ ID NO: 84 is the determined DNA sequence for the *C. trachomatis* LGV II clone 19-A5-54, sharing homology to the cryptic plasmid gene.

15 SEQ ID NO: 85 is the determined DNA sequence for the C. trachomatis LGV II clone 17-E11-72, sharing partial homology to the OppC 2 and pmpD genes.

SEQ ID NO: 86 is the determined DNA sequence for the *C. trachomatis* LGV II clone 17-C1-77, sharing partial homology to the CT857 and CT858 open reading frames.

SEQ ID NO: 87 is the determined DNA sequence for the *C. trachomatis* LGV II clone 15-H2-76, sharing partial homology to the pmpD and SycE genes, and to the CT089 ORF.

SEQ ID NO: 88 is the determined DNA sequence for the *C. trachomatis* LGV II clone 15-A3-26, sharing homology to the CT858 ORF.

25 SEQ ID NO: 89 is the determined amino acid sequence for the C. pnuemoniae clone Cp SWIB-His.

SEQ ID NO: 90 is the determined amino acid sequence for the C. trachomatis LGV II clone CtL2 LPDA FL.

SEQ ID NO: 91 is the determined amino acid sequence for the C.

30 pnuemoniae clone CpS13-His.

SEQ ID NO: 92 is the determined amino acid sequence for the C. trachomatis LGV II clone CtL2 TSA FL.

SEQ ID NO: 93 is the amino acid sequence for Ct-Swib 43-61 peptide from C. trachomatis LGV II.

SEQ ID NO: 94 is the amino acid sequence for Ct-Swib 48-67 peptide from C. trachomatis LGV II.

5 SEQ ID NO: 95 is the amino acid sequence for Ct-Swib 52-71 peptide from C. trachomatis LGV II.

SEQ ID NO: 96 is the amino acid sequence for Ct-Swib 58-77 peptide from C. trachomatis LGV II.

SEQ ID NO: 97 is the amino acid sequence for Ct-Swib 63-82 peptide from C. trachomatis LGV II.

SEQ ID NO: 98 is the amino acid sequence for Ct-Swib 51-66 peptide from C. trachomatis LGV II.

SEQ ID NO: 99 is the amino acid sequence for Cp-Swib 52-67 peptide from *C. pneumonia*.

15 SEQ ID NO: 100 is the amino acid sequence for Cp-Swib 37-51 peptide from C. pneumonia.

SEQ ID NO: 101 is the amino acid sequence for Cp-Swib 32-51 peptide from $\it C. pneumonia$.

SEQ ID NO: 102 is the amino acid sequence for Cp-Swib 37-56 peptide from C. meumonia.

SEQ ID NO: 103 is the amino acid sequence for Ct-Swib 36-50 peptide from C. trachomatis.

SEQ ID NO: 104 is the amino acid sequence for Ct-S13 46-65 peptide from C. trachomatis.

25 SEQ ID NO: 105 is the amino acid sequence for Ct-S13 60-80 peptide from C. trachomatis.

SEQ ID NO: 106 is the amino acid sequence for Ct-S13 1-20 peptide from C. trachomatis.

SEQ ID NO: 107 is the amino acid sequence for Ct-S13 46-65 peptide from C. trachomatis.

SEQ ID NO: 108 is the amino acid sequence for Ct-S13 56-75 peptide from C. trachomatis.

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SEQ ID NO: 109 is the amino acid sequence for Cp-S13 56-75 peptide from C. pneumoniae.

SEQ ID NO: 110 is the determined DNA sequence for the *C. trachomatis* LGV II clone 21-G12-60, containing partial open reading frames for 5 hypothetical proteins CT875. CT229 and CT228.

SEQ ID NO: 111 is the determined DNA sequence for the *C. trachomatis* LGV II clone 22-B3-53, sharing homology to the CT110 ORF of GroEL.

SEQ ID NO: 112 is the determined DNA sequence for the $\it C$. trachomatis LGV II clone 22-A1-49, sharing partial homology to the CT660 and CT659 ORFs.

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SEQ ID NO: 113 is the determined DNA sequence for the *C. trachomatis* LGV II clone 17-E2-9, sharing partial homology to the CT611 and CT 610 ORFs.

SEQ ID NO: 114 is the determined DNA sequence for the C.

15 trachomatis LGV II clone 17-C10-31, sharing partial homology to the CT858 ORF.

SEQ ID NO: 115 is the determined DNA sequence for the $\it C.$ trachomatis LGV II clone 21-C7-8, sharing homology to the dnaK-like gene.

SEQ ID NO: 116 is the determined DNA sequence for the *C. trachomatis* LGV II clone 20-G3-45, containing part of the pmpB gene CT413.

20 SEQ ID NO: 117 is the determined DNA sequence for the C. trachomatis LGV II clone 18-C5-2, sharing homology to the S1 ribosomal protein ORF.

SEQ ID NO: 118 is the determined DNA sequence for the $\it C$. trachomatis LGV II clone 17-C5-19, containing part of the ORFs for CT431 and CT430.

SEQ ID NO: 119 is the determined DNA sequence for the C. trachomatis LGV II clone 16-D4-22, contains partial sequences of ORF3 and ORF4 of the plasmid for growth within mammalian cells.

SEQ ID NO: 120 is the determined full-length DNA sequence for the $\it C.$ trachomatis serovar LGV II Cap1 gene CT529.

30 SEQ ID NO: 121 is the predicted full-length amino acid sequence for the C. trachomatis serovar LGV II Cap1 gene CT529.

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SEQ ID NO: 122 is the determined full-length DNA sequence for the *C. trachomatis* serovar E Cap1 gene CT529.

SEQ ID NO: 123 is the predicted full-length amino acid sequence for the C. trachomatis serovar E Cap1 gene CT529.

5 SEQ ID NO: 124 is the determined full-length DNA sequence for the C. trachomatis serovar 1A Cap1 gene CT529.

SEQ ID NO: 125 is the predicted full-length amino acid sequence for the C. trachomatis serovar 1A Cap1 gene CT529.

SEQ ID NO: 126 is the determined full-length DNA sequence for the C.

trachomatis serovar G Cap1 gene CT529.

SEQ ID NO: 127 is the predicted full-length amino acid sequence for the C. trachomatis serovar G Cap1 gene CT529.

SEQ ID NO: 128 is the determined full-length DNA sequence for the *C. trachomatis* serovar F1 NII Cap1 gene CT529.

15 SEQ ID NO: 129 is the predicted full-length amino acid sequence for the C. trachomatis serovar F1 NII Cap1 gene CT529.

SEQ ID NO: 130 is the determined full-length DNA sequence for the $\it C$ -trachomatis serovar L1 Cap1 gene CT529.

SEQ ID NO: 131 is the predicted full-length amino acid sequence for the

C. trachomatis serovar L1 Cap1 gene CT529.

SEQ ID NO: 132 is the determined full-length DNA sequence for the C. trachomatis serovar L3 Cap1 gene CT529.

SEQ ID NO: 133 is the predicted full-length amino acid sequence for the C. trachomatis serovar L3 Cap1 gene CT529.

SEQ ID NO: 134 is the determined full-length DNA sequence for the C. trachomatis servora Ba Cap1 gene CT529.

SEQ ID NO: 135 is the predicted full-length amino acid sequence for the C. trachomatis serovar Ba Cap1 gene CT529.

SEQ ID NO: 136 is the determined full-length DNA sequence for the C. trachomatis serovar MOPN Cap1 gene CT529.

SEQ ID NO: 137 is the predicted full-length amino acid sequence for the C. trachomatis serovar MOPN Cap1 gene CT529.

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SEQ ID NO: 138 is the determined amino acid sequence for the Cap1

CT529 ORF peptide #124-139 of C. trachomatis serovar L2.

SEQ ID NO: 139 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #132-147 of *C. trachomatis* serovar L2.

5 SEQ ID NO: 140 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #138-155 of C. trachomatis serovar L2.

SEQ ID NO: 141 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #146-163 of *C. trachomatis* serovar L2.

SEQ ID NO: 142 is the determined amino acid sequence for the Cap1

10 CT529 ORF peptide #154-171 of C. trachomatis serovar L2.

SEQ ID NO: 143 is the determined amino acid sequence for the Cap1 CT529 ORF pentide #162-178 of *C. trachomatis* serovar L2.

SEQ ID NO: 144 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #138-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 145 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #139-147 of *C. trachomatis* serovar L2.

SEQ ID NO: 146 is the determined amino acid sequence for the Cap1 CT529 ORF peptide #140-147 of *C. trachomatis* seroyar L2.

SEO ID NO: 147 is the determined amino acid sequence for the Cap1

20 CT529 ORF peptide #138-146 of C. trachomatis serovar L2.

SEQ ID NO: 148 is the determined amino acid sequence for the Cap1

CT529 ORF peptide #138-145 of C. trachomatis serovar L2.

SEQ ID NO: 149 is the determined amino acid sequence for the Cap1

CT529 ORF peptide # F140->I of C. trachomatis serovar L2.

SEQ ID NO: 150 is the determined amino acid sequence for the Cap1 CT529 ORF peptide ##S139>Ga of *C. trachomatis* serovar L2.

SEQ ID NO: 151 is the determined amino acid sequence for the Cap1

CT529 ORF peptide ##S139>Gb of C. trachomatis serovar L2.

SEQ ID NO: 152 is the determined amino acid sequence for the peptide

2 C7.8-6 of the 216aa ORF of C. trachomatis serovar L2.

SEQ ID NO: 153 is the determined amino acid sequence for the peptide # 2 C7.8-7 of the 216aa ORF of *C. trachomatis* serovar L2.

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SEQ ID NO: 154 is the determined amino acid sequence for the peptide #2 C7.8-8 of the 216aa ORF of *C. trachomatis* serovar L2.

SEQ ID NO: 155 is the determined amino acid sequence for the peptide #2 C7.8-9 of the 216aa ORF of *C. trachomatis* serovar L2.

SEQ ID NO: 156 is the determined amino acid sequence for the peptide # 2 C7.8-10 of the 216aa ORF of C. trachomatis serovar L2.

SEQ ID NO: 157 is the determined amino acid sequence for the 53 amino acid residue peptide of the 216aa ORF within clone 2C7.8 of *C. trachomatis* server 1.2

SEQ ID NO: 158 is the determined amino acid sequence for the 52 amino acid residue peptide of the CT529 ORF within clone 2C7.8 of C. trachomatis serovar L2.

SEQ ID NO: 159 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 serovar L2.

15 SEQ ID NO: 160 is the determined DNA sequence for the 5' (reverse) primer for cloning full-length CT529 serovar L2.

SEQ ID NO: 161 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 for serovars other than L2 and MOPN.

SEQ ID NO: 162 is the determined DNA sequence for the 5' (reverse)
20 primer for cloning full-length CT529 serovars other than L2 and MOPN.

SEQ ID NO: 163 is the determined DNA sequence for the 5' (forward) primer for cloning full-length CT529 serovar MOPN.

SEQ ID NO: 164 is the determined DNA sequence for the 5' (reverse) primer for cloning full-length CT529 serovar MOPN.

25 SEQ ID NO: 165 is the determined DNA sequence for the 5' (forward) primer for pBIB-KS.

SEQ ID NO: 166 is the determined DNA sequence for the 5' (reverse) primer for pBIB-KS.

SEQ ID NO: 167 is the determined amino acid sequence for the 9-mer openitope pentide Cap1#139-147 from serovar L2.

SEQ ID NO: 168 is the determined amino acid sequence for the 9-mer epitope peptide Cap1#139-147 from serovar D.

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SEQ ID NO: 169 is the determined full-length DNA sequence for the C. trachomatis pmpI (CT874) gene.

SEQ ID NO: 170 is the determined full-length DNA sequence for the $\it C.$ $\it trachomatis pmpG$ gene.

5 SEQ ID NO: 171 is the determined full-length DNA sequence for the C. trachomatis pmpE gene.

SEQ ID NO: 172 is the determined full-length DNA sequence for the $\it C.$ trachomatis pmpD gene.

SEQ ID NO: 173 is the determined full-length DNA sequence for the $\it C.$ trachomatis pmpC gene.

SEQ ID NO: 174 is the determined full-length DNA sequence for the $\it C.$ trachomatis pmpB gene.

SEQ ID NO: 175 is the predicted full-length amino acid sequence for the C. trachomatis pmpI gene.

15 SEQ ID NO: 176 is the predicted full-length amino acid sequence for the C. trachomatis pmpG gene.

SEQ ID NO: 177 is the predicted full-length amino acid sequence for the C. trachomatis pmpE gene.

SEQ ID NO: 178 is the predicted full-length amino acid sequence for the C. trachomatis pmpD gene.

SEQ ID NO: 179 is the predicted full-length amino acid sequence for the $\it C. trachomatis pmpC$ gene.

SEQ ID NO: 180 is the predicted full-length amino acid sequence for the $\it C.\ trachomatis$ pmpB gene.

25 SEQ ID NO: 181 is the determined DNA sequence minus the signal sequence for the C. trachomatis pmpI gene.

SEQ ID NO: 182 is a subsequently determined full-length DNA sequence for the $\it C. trachomatis \ pmpG \ gene.$

SEQ ID NO: 183 is the determined DNA sequence minus the signal $_{30}$ sequence for the C. trachomatis pmpE gene.

SEQ ID NO: 184 is a first determined DNA sequence representing the . carboxy terminus for the C. trachomatis pmpD gene.

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SEQ ID NO: 185 is a second determined DNA sequence representing the amino terminus minus the signal sequence for the *C. trachomatis* pmpD gene.

SEQ ID NO: 186 is a first determined DNA sequence representing the carboxy terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 187 is a second determined DNA sequence representing the amino terminus minus the signal sequence for the C. trachomatis pmpC gene.

SEQ ID NO: 188 is the determined DNA sequence representing the C. pneumoniae serovar MOMPS pmp gene in a fusion molecule with Ra12.

SEQ ID NO: 189 is the predicted amino acid sequence minus the signal sequence for the C. trachomatis pmpI gene.

SEQ ID NO: 190 is subsequently predicted amino acid sequence for the C. trachomatis pmpG gene.

SEQ ID NO: 191 is the predicted amino acid sequence minus the signal sequence for the C. trachomatis pmpE gene.

15 SEQ ID NO: 192 is a first predicted amino acid sequence representing the carboxy terminus for the C. trachomatis pmpD gene.

SEQ ID NO: 193 is a second predicted amino acid sequence representing the Amino terminus minus the signal sequence for the *C. trachomatis* pmpD gene.

SEQ ID NO: 194 is a first predicted amino acid sequence representing
the Carboxy terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 195 is a second predicted amino acid sequence representing the Amino terminus for the *C. trachomatis* pmpC gene.

SEQ ID NO: 196 is the predicted amino acid sequence representing the C. pneumoniae scroyar MOMPS pmp gene in a fusion molecule with Ral2.

SEQ ID NO: 197 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 198 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 199 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpC gene in the SKB vaccine vector.

SEQ ID NO: 200 is the determined DNA sequence for the 5° oligo primer for cloning the C. trachomatis pmpD gene in the SKB vaccine vector.

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SEQ ID NO: 201 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpD gene in the SKB vaccine vector.

SEQ ID NO: 202 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpD gene in the SKB vaccine vector.

SEQ ID NO: 203 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpE gene in the SKB vaccine vector.

SEQ ID NO: 204 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpE gene in the SKB vaccine vector.

SEQ ID NO: 205 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 206 is the determined DNA sequence for the 3' oligo primer for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 207 is the determined DNA sequence for the 5° oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpC gene in the 15 pET17b vector.

SEQ ID NO: 208 is the determined DNA sequence for the 3° oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 209 is the determined DNA sequence for the 5' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 210 is the determined DNA sequence for the 3' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpC gene in the pET17b vector.

SEQ ID NO: 211 is the determined DNA sequence for the 5° oligo primer for cloning the amino terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 213 is the determined DNA sequence for the 5' oligo primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpD gene in the pET17b vector.

SEQ ID NO: 214 is the determined DNA sequence for the 3' oligo
primer for cloning the carboxy terminus portion of the *C. trachomatis* pmpD gene in the
pET17b vector.

SEQ ID NO: 215 is the determined DNA sequence for the 5' oligo primer for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

SEQ ID NO: 216 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpE gene in the pET17b vector.

SEQ ID NO: 217 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

SEQ ID NO: 218 is the amino acid sequence for the insertion sequence for cloning the *C. trachomatis* pmpE gene in the pET17b vector.

15 SEQ ID NO: 219 is the determined DNA sequence for the 5' oligo primer for cloning the C. trachomatis pmpG gene in the pET17b vector.

SEQ ID NO: 220 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpG gene in the pET17b vector.

SEQ ID NO: 221 is the amino acid sequence for the insertion sequence

20 for cloning the *C. trachomatis* pmpG gene in the pET17b vector.

SEQ ID NO: 222 is the determined DNA sequence for the 5' oligo primer for cloning the C. trachomatis pmpI gene in the pET17b vector.

SEQ ID NO: 223 is the determined DNA sequence for the 3' oligo primer for cloning the C. trachomatis pmpI gene in the pET17b vector.

SEQ ID NO: 224 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 1-20.

SEQ ID NO: 225 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 6-25.

SEQ ID NO: 226 is the determined amino acid sequence for the C.

pneumoniae Swib peptide 12-31.

SEQ ID NO: 227 is the determined amino acid sequence for the $\it C.$ pneumoniae Swib peptide 17-36.

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SEQ ID NO: 228 is the determined amino acid sequence for the *C. pneumoniae* Swib pertide 22-41.

SEQ ID NO: 229 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 27-46.

5 SEQ ID NO: 230 is the determined amino acid sequence for the C. pneumoniae Swib pertide 42-61.

SEQ ID NO: 231 is the determined amino acid sequence for the *C. pneumoniae* Swib pertide 46-65.

SEQ ID NO: 232 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 51-70.

SEQ ID NO: 233 is the determined amino acid sequence for the *C. pneumoniae* Swib peptide 56-75.

SEQ ID NO: 234 is the determined amino acid sequence for the Cpneumoniae Swib peptide 61-80.

15 SEQ ID NO: 235 is the determined amino acid sequence for the C. pneumoniae Swib pertide 66-87.

SEQ ID NO: 236 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 103-122.

SEQ ID NO: 237 is the determined amino acid sequence for the C.
trachomatis OMCB peptide 108-127.

SEQ ID NO: 238 is the determined amino acid sequence for the C. trachomatis OMCB peptide 113-132.

SEQ ID NO: 239 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 118-137.

SEQ ID NO: 240 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 123-143.

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SEQ ID NO: 241 is the determined amino acid sequence for the C. trachomatis OMCB peptide 128-147,

SEQ ID NO: 242 is the determined amino acid sequence for the C. trachomatis OMCB peptide 133-152.

SEQ ID NO: 243 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 137-156,

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SEQ ID NO: 244 is the determined amino acid sequence for the C. trachomatis OMCB pentide 142-161.

SEQ ID NO: 245 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 147-166.

5 SEQ ID NO: 246 is the determined amino acid sequence for the C. trachomatis OMCB peptide 152-171.

SEQ ID NO: 247 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 157-176.

SEQ ID NO: 248 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 162-181.

SEQ ID NO: 249 is the determined amino acid sequence for the *C. trachomatis* OMCB pentide 167-186.

SEQ ID NO: 250 is the determined amino acid sequence for the *C. trachomatis* OMCB peptide 171-190.

15 SEQ ID NO: 251 is the determined amino acid sequence for the C. trachomatis OMCB pentide 171-186.

SEQ ID NO: 252 is the determined amino acid sequence for the $\it C.$ trachomatis OMCB peptide 175-186.

SEQ ID NO: 252 is the determined amino acid sequence for the C.

20 trachomatis OMCB peptide 175-186.

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SEQ ID NO: 253 is the determined amino acid sequence for the *C. pneumoniae* OMCB peptide 185-198.

SEQ ID NO: 254 is the determined amino acid sequence for the $\it C. trachomatis$ TSA peptide 96-115.

SEQ ID NO: 255 is the determined amino acid sequence for the C. trachomatis TSA peptide 101-120.

SEQ ID NO: 256 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 106-125.

SEQ ID NO: 257 is the determined amino acid sequence for the C. 30 trachomatis TSA pertide 111-130.

SEQ ID NO: 258 is the determined amino acid sequence for the C. trachomatis TSA peptide 116-135.

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SEQ ID NO: 259 is the determined amino acid sequence for the *C. trachomatis* TSA pentide 121-140.

SEQ ID NO: 260 is the determined amino acid sequence for the C. trachomatis TSA peptide 126-145.

SEQ ID NO: 261 is the determined amino acid sequence for the *C. trachomatis* TSA peptide 131-150.

SEQ ID NO: 262 is the determined amino acid sequence for the $\it C.trachomatis$ TSA peptide 136-155.

SEQ ID NO: 263 is the determined full-length DNA sequence for the C.

trachomatis CT529/Cap 1 gene serovar I.

SEQ ID NO: 264 is the predicted full-length amino sequence for the C. trachomatis CT529/Cap 1 gene serovar I.

SEQ ID NO: 265 is the determined full-length DNA sequence for the C. trachomatis CT529/Cap 1 gene serovar K.

SEQ ID NO: 266 is the predicted full-length amino sequence for the C. trachomatis CT529/Cap 1 gene serovar K.

SEQ ID NO: 267 is the determined DNA sequence for the $\it C.$ trachomatis clone 17-G4-36 sharing homology to part of the ORF of DNA-dirrected RNA polymerase beta subunit- CT315 in serD.

SEQ ID NO: 268 is the determined DNA sequence for the partial sequence of the C. trachomatis CT016 gene in clone 2E10.

SEQ ID NO: 269 is the determined DNA sequence for the partial sequence of the C. trachomatis tRNA syntase gene in clone 2E10.

SEQ ID NO: 270 is the determined DNA sequence for the partial sequence for the *C. trachomatis* clpX gene in clone 2E10.

SEQ ID NO: 271 is a first determined DNA sequence for the *C. trachomatis* clone CtL2gam-30 representing the 5'end.

SEQ ID NO: 272 is a second determined DNA sequence for the C. trachomatis clone CtL2gam-30 representing the 3'end.

SEQ ID NO: 273 is the determined DNA sequence for the C. trachomatis clone CtL2gam-28.

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- SEQ ID NO: 274 is the determined DNA sequence for the C. trachomatis clone CtL2gam-27.
- SEQ ID NO: 275 is the determined DNA sequence for the C. trachomatis clone CtL2gam-26.
- 5 SEQ ID NO: 276 is the determined DNA sequence for the C. trachomatis clone CtL2pam-24.
 - SEQ ID NO: 277 is the determined DNA sequence for the C. trachomatis clone CtL2gam-23.
- SEQ ID NO: 278 is the determined DNA sequence for the C.

 10 trachomatis clone CtL2gam-21.
 - SEQ ID NO: 279 is the determined DNA sequence for the *C. trachomatis* clone CtL2eam-18.
 - SEQ ID NO: 280 is the determined DNA sequence for the $\it C.$ trachomatis clone CtL2gam-17.
- 15 SEQ ID NO: 281 is a first determined DNA sequence for the C. trachomatis clone CtL2gam-15 representing the 5' end.
 - SEQ ID NO: 282 is a second determined DNA sequence for the C. trachomatis clone CtL2gam-15 representing the 3' end.
- SEQ ID NO: 283 is the determined DNA sequence for the C. 20 trachomatis clone CtL2gam-13.
 - SEQ ID NO: 284 is the determined DNA sequence for the $\it C.$ trachomatis clone CtL2gam-10.
 - SEQ ID NO: 285 is the determined DNA sequence for the $\it C.$ trachomatis clone CtL2gam-8.
 - SEQ ID NO: 286 is a first determined DNA sequence for the C. trachomatis clone CtL2gam-6 representing the 5' end.
 - SEQ ID NO: 287 is a second determined DNA sequence for the C. trachomatis clone CtL2gam-6 representing the 3' end.
- SEQ ID NO: 288 is the determined DNA sequence for the C.

 30 trachomatis clone CtL2gam-5.
 - SEQ ID NO: 289 is the determined DNA sequence for the $\it C.$ trachomatis clone CtL2gam-2.

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SEQ ID NO: 290 is the determined DNA sequence for the C. trachomatis clone CtL2gam-1.

SEQ ID NO: 291 is the determined full-length DNA sequence for the C. pneumoniae homologue of the CT529 gene.

5 SEQ ID NO: 292 is the predicted full-length amino acid sequence for the C. pneumoniae homologue of the CT529 gene.

SEQ ID NO: 293 is the determined DNA sequence for the insertion sequence for cloning the *C. trachomatis* pmpG gene in the SKB vaccine vector.

SEQ ID NO: 294 is the amino acid sequence of an open reading frame of clone CT603.

SEQ ID NO: 295 is the amino acid sequence of a first open reading frame of clone CT875.

SEQ ID NO: 296 is the amino acid sequence of a second open reading frame of clone CT875.

15 SEQ ID NO: 297 is the amino acid sequence of a first open reading frame of clone CT858.

SEQ ID NO: 298 is the amino acid sequence of a second open reading frame of clone CT858.

SEQ ID NO: 299 is the amino acid sequence of an open reading frame of clone CT622.

 $\,$ SEQ ID NO: 300 is the amino acid sequence of an open reading frame of clone CT610.

SEQ ID NO: 301 is the amino acid sequence of an open reading frame of clone CT396.

25 SEQ ID NO: 302 is the amino acid sequence of an open reading frame of clone CT318.

SEQ ID NO: 304 is the amino acid sequence for *C. trachomatis*, serovar L2 rCt529c1-125 having a modified N-terminal sequence (6-His tag).

SEQ ID NO: 305 is the amino acid sequence for *C. trachomatis*, serovar L2 rCt529c1-125.

SEQ ID NO: 306 is the sense primer used in the synthesis of the $\mbox{PmpA(N-term)}$ fusion protein.

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\$25\$ SEQ ID NO: 307 is the antisense primer used in the synthesis of the $$\operatorname{PmpA(N-term)}$$ fusion protein.

SEQ ID NO: 308 is the DNA sequence encoding the PmpA(N-term) fusion protein.

5 SEQ ID NO: 309 is the amino acid sequence of the PmpA(N-term)

fusion protein.

SEQ ID NO: 310 is the sense primer used in the synthesis of the $\mbox{PmpA}(\mbox{C-term})$ fusion protein.

SEQ ID NO: 311 is the antisense primer used in the synthesis of the
10 PmpA(C-term) fusion protein.

SEQ ID NO: 312 is the DNA sequence encoding the PmpA(C-term) fusion protein.

SEQ ID NO: 313 is the amino acid sequence of the PmpA(C-term) fusion protein.

15 SEQ ID NO: 314 is the sense primer used in the synthesis of the PmpF(N-term) fusion protein.

SEQ ID NO: 315 is the antisense primer used in the synthesis of the PmpF(N-term) fusion protein.

SEQ ID NO: 316 is the DNA sequence encoding the PmpF(N-term) 20 fusion protein.

SEQ ID NO: 317 is the amino acid sequence of the PmpF(N-term) fusion protein.

SEQ ID NO: 318 is the sense primer used in the synthesis of the PmpF(C-term) fusion protein.

SEQ ID NO: 319 is the antisense primer used in the synthesis of the PmpF(C-term) fusion protein.

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SEQ ID NO: 320 is the DNA sequence encoding the PmpF(C-term) fusion protein.

SEQ ID NO: 321 is the amino acid sequence of the PmpF(C-term) fusion protein.

SEQ ID NO: 322 is the sense primer used in the synthesis of the PmpH (CT412) (N-term) fusion protein.

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SEQ ID NO: 323 is the antisense primer used in the synthesis of the PmpH(N-term) fusion protein.

SEQ ID NO: 324 is the DNA sequence encoding the PmpH(N-term) fusion protein.

5 SEO ID NO: 325 is the amino acid sequence of the PmpH(N-term) fusion protein.

SEQ ID NO: 326 is the sense primer used in the synthesis of the PmpH(C-term) fusion protein.

SEO ID NO: 327 is the antisense primer used in the synthesis of the PmpH(C-term) fusion protein.

SEO ID NO: 328 is the DNA sequence encoding the PmpH(C-term) fusion protein.

SEQ ID NO: 329 is the amino acid sequence of the PmpH(C-term) fusion protein.

SEO ID NO: 330 is the sense primer used in the synthesis of the PmpB(1) fusion protein.

SEQ ID NO: 331 is the antisense primer used in the synthesis of the PmpB(1) fusion protein.

SEO ID NO: 332 is the DNA sequence encoding the PmpB(1) fusion 20 protein.

SEQ ID NO: 333 is the amino acid sequence of the PmpB(1) fusion protein.

SEO ID NO: 334 is the sense primer used in the synthesis of the PmpB(2) fusion protein.

SEQ ID NO: 335 is the antisense primer used in the synthesis of the 25 PmpB(2) fusion protein.

SEQ ID NO: 336 is the DNA sequence encoding the PmpB(2) fusion protein.

SEQ ID NO: 337 is the amino acid sequence of the PmpB(2) fusion

SEO ID NO: 338 is the sense primer used in the synthesis of the PmpB(3) fusion protein.

20 protein.

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SEQ ID NO: 339 is the antisense primer used in the synthesis of the PmpB(3) fusion protein.

SEQ ID NO: 340 is the DNA sequence encoding the PmpB(3) fusion protein.

SEQ ID NO: 341 is the amino acid sequence of the PmpB(3) fusion protein.

SEQ ID NO: 342 is the sense primer used in the synthesis of the PmpB(4) fusion protein.

SEQ ID NO: 343 is the antisense primer used in the synthesis of the PmpB(4) fusion protein.

SEQ ID NO: 344 is the DNA sequence encoding the PmpB(4) fusion protein.

SEQ ID NO: 345 is the amino acid sequence of the PmpB(4) fusion protein.

15 SEQ ID NO: 346 is the sense primer used in the synthesis of the PmpC(1) fusion protein.

SEQ ID NO: 347 is the antisense primer used in the synthesis of the PmpC(1) fusion protein.

SEQ ID NO: 348 is the DNA sequence encoding the PmpC(1) fusion

SEQ ID NO: 349 is the amino acid sequence of the PmpC(1) fusion protein.

SEQ ID NO: 350 is the sense primer used in the synthesis of the PmpC(2) fusion protein.

SEQ ID NO: 351 is the antisense primer used in the synthesis of the PmpC(2) fusion protein.

SEQ ID NO: 352 is the DNA sequence encoding the PmpC(2) fusion protein.

SEQ ID NO: 353 is the amino acid sequence of the PmpC(2) fusion protein.

SEQ ID NO: 354 is the sense primer used in the synthesis of the PmpC(3) fusion protein.

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SEQ ID NO: 355 is the antisense primer used in the synthesis of the PmpC(3) fusion protein.

SEQ ID NO: 356 is the DNA sequence encoding the PmpC(3) fusion protein.

SEQ ID NO: 357 is the amino acid sequence of the PmpC(3) fusion protein.

SEQ ID NO: 358 is the DNA sequence of the oppA1 protein, devoid of the first trans-membrane domain.

SEQ ID NO: 359 is the full length DNA sequence of CT139.

SEQ ID NO: 360 is the full length DNA sequence of ORF-3.

SEQ ID NO: 361 is the full length DNA sequence of CT611.

SEQ ID NO: 362 is the amino acid sequence of oppA1 starting from amino acid 22.

SEQ ID NO: 363 is the amino acid sequence of CT139.

SEO ID NO: 364 is the amino acid sequence of ORF-3.

SEQ ID NO: 365 is the amino acid sequence of CT611.

SEQ ID NO: 366 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0275, of the Chlamydia trachomatis gene CT190.

SEQ ID NO: 367 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0407, of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 368 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0720, of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 369 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0716, of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 370 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0519, of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 371 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0520, of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 372 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0078, of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 373 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0628, of the Chlamydia trachomatis gene CT509.

SEQ ID NO: 374 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0540, of the Chlamydia trachomatis gene CT414.

SEQ ID NO: 375 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, pmp20, of the Chlamydia trachomatis gene CT413.

SEQ ID NO: 376 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0081, of the Chlamydia trachomatis gene CT315.

SEQ ID NO: 377 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0761, of the Chlamydia trachomatis gene CT610.

SEQ ID NO: 378 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0557, of the Chlamydia trachomatis gene CT443.

SEQ ID NO: 379 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0833, of the Chlamydia trachomatis gene CT557.

SEQ ID NO: 380 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0134, of the Chlamydia trachomatis gene CT604.

SEQ ID NO: 381 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0388, of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 382 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn1028, of the Chlamydia trachomatis gene CT376.

SEQ ID NO: 383 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0875, of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 384 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0908, of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 385 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0728, of the Chlamydia trachomatis gene CT622.

SEQ ID NO: 386 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0275, of the Chlamydia trachomatis gene CT190.

SEQ ID NO: 387 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0407, of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 388 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0720, of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 389 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0716, of the Chlamydia trachomatis gene CT660.

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SEQ ID NO: 390 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0519, of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 391 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0520, of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 392 sets forth the amino acid sequence for the Chlamydia

pneumoniae homologue, CPn0078, of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 393 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0628, of the Chlamydia trachomatis gene CT509.

SEQ ID NO: 394 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0540, of the Chlamydia trachomatis gene CT414.

SEQ ID NO: 395 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, pmp20, of the Chlamydia trachomatis gene CT413.

SEQ ID NO: 396 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0081, of the Chlamydia trachomatis gene CT315.

SEQ ID NO: 397 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0761, of the Chlamydia trachomatis gene CT610.

pneumoniae homologue, CPn0557, of the Chlamydia trachomatis gene CT443.

SEQ ID NO: 399 sets forth the amino acid sequence for the Chlamydia

SEQ ID NO: 398 sets forth the amino acid sequence for the Chlamydia

pneumoniae homologue, CPn0833, of the Chlamydia trachomatis gene CT557.

SEQ ID NO: 400 sets forth the amino acid sequence for the Chlamydia

pneumoniae homologue, CPn0134, of the Chlamydia trachomatis gene CT604.
SEQ ID NO: 401 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0388, of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 402 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue. CPn1028, of the Chlamydia trachomatis gene CT376.

SEQ ID NO: 403 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0875, of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 404 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0908, of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 405 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0728, of the Chlamydia trachomatis gene CT622.

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SEQ ID NO: 406 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT287.

SEQ ID NO: 407 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT858.

SEQ ID NO: 408 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 409 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 410 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 411 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 412 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT622.

SEQ ID NO: 413 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT610.

SEQ ID NO: 414 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT604.

SEQ ID NO: 415 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT557.

SEQ ID NO: 416 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT509.

SEQ ID NO: 417 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT443.

SEQ ID NO: 418 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT431.

SEQ ID NO: 419 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT430.

SEQ ID NO: 420 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT414.

SEQ ID NO: 421 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT413.

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SEQ ID NO: 422 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT396.

SEQ ID NO: 423 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT376.

SEQ ID NO: 424 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT318.

SEQ ID NO: 425 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT315.

SEQ ID NO: 426 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT104.

SEQ ID NO: 427 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT103.

SEQ ID NO: 428 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 429 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 430 sets forth the full-length serovar D DNA sequence of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 431 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT858.

SEQ ID NO: 432 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT764.

SEQ ID NO: 433 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT734.

SEQ ID NO: 434 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT660.

SEQ ID NO: 435 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT659.

SEQ ID NO: 436 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT622.

SEQ ID NO: 437 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT610.

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- SEQ ID NO: 438 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT604.
- SEQ ID NO: 439 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT557.
- SEQ ID NO: 440 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT509.
- SEQ ID NO: 441 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT443.
- SEQ ID NO: 442 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT431.
- SEQ ID NO: 443 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT430.
- SEQ ID NO: 444 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT414.
- SEQ ID NO: 445 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT413.
- SEQ ID NO: 446 sets forth the full-length serovar D amino acid sequence of the Chlamydia trachomatis gene CT396.
- SEQ ID NO: 447 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT376.
- SEQ ID NO: 448 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT318.
- SEQ ID NO: 449 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT315.
- SEQ ID NO: 450 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT104.
- SEQ ID NO: 451 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT103.
- SEQ ID NO: 452 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT102.
- SEQ ID NO: 453 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT098.

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SEQ ID NO: 454 sets forth the full length serovar D amino acid sequence of the Chlamydia trachomatis gene CT042.

SEQ ID NO: 455 corresponds to the DNA sequence of CPn0894, which is the CP homologue of CT751 (amn), which was identified in clones CTL2-1, and CTL2-5.

SEQ ID NO: 456 corresponds to the DNA sequence of CPn0074, which is the CP homologue of CT322 (tuf), which was identified in clone CTL2-2.

SEQ ID NO: 457 corresponds to the DNA sequence of CPn0122, which is the CP homologue of CT032 (metG), which was identified in clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 458 corresponds to the DNA sequence of CPn0121, which is the CP homologue of CT031, which was identified in clone CTL2-3(5')(3').

SEQ ID NO: 459 corresponds to the DNA sequence of CPn0120, which is the CP homologue of CT030 (gmK), which was identified in clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 460 corresponds to the DNA sequence of CPn0359, which is the CP homologue of CT064 (lepA), which was identified in clone CTL2gam5.

SEQ ID NO: 461 corresponds to the DNA sequence of CPn0414, which is the CP homologue of CT265 (accA), which was identified in clone CTL2-6.

SEQ ID NO: 462 corresponds to the DNA sequence of CPn0413, which is the CP homologue of CT264 (msbA), which was identified in clone CTL2-6.

SEQ ID NO: 463 corresponds to the DNA sequence of CPn0394, which is the CP homologue of CT256 which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 464 corresponds to the DNA sequence of CPn0395, which is the CP homologue of CT257 which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 465 corresponds to the DNA sequence of CPn0487, which is the CP homologue of CT384 which was identified in clones CTL2gam6(3') and CTL2-11(3').

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SEQ ID NO: 466 corresponds to the DNA sequence of CPn0592, which is the CP homologue of CT473, which was identified in clone CTL2-8b.

SEQ ID NO: 467 corresponds to the DNA sequence of CPn0593, which is the CP homologue of CT474, which was identified in clone CTL2-8b.

SEQ ID NO: 468 corresponds to the DNA sequence of CPn0197, which is the CP homologue of CT139 (oppA1), which was identified in clone CTL2-8b.

SEQ ID NO: 469 corresponds to the DNA sequence of CPn0363, which is the CP homologue of CT060 (flhA), which was identified in clone CTL2-8b.

SEQ ID NO: 470 corresponds to the DNA sequence of CPn0301, which is the CP homologue of CT242, which was identified in clone CTL2gam8.

SEQ ID NO: 471 corresponds to the DNA sequence of CPn0302, which is the CP homologue of CT243 (lpxD), which was identified in clone CTL2gam8.

SEQ ID NO: 472 corresponds to the DNA sequence of CPn0324, which is the CP homologue of CT089 (lcrE), which was identified in clones CTL2-9, CTL2gam1, CTL2gam17 and CTL2-19(5').

SEQ ID NO: 473 corresponds to the DNA sequence of CPn0761, which is the CP homologue of CT610, which was identified in clone CTL2-10(5')(3').

SEQ ID NO: 474 corresponds to the DNA sequence of CPn0760, which is the CP homologue of CT611, which was identified in clone CTL2-10(5').

SEQ ID NO: 475 corresponds to the DNA sequence of CPn0329, which is the CP homologue of CT154, which was identified in clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 476 corresponds to the DNA sequence of CPn0990, which is the CP homologue of CT833 (infC), which was identified in clone CTL2-12.

SEQ ID NO: 477 corresponds to the DNA sequence of CPn0984, which is the CP homologue of CT827 (nrdA), which was identified in clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 478 corresponds to the DNA sequence of CPn0985 which is the CP homologue of CT828 (nrdB) which was identified in clones CTL2-16(3') CTL2gam15(3').

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SEQ ID NO: 479 corresponds to the DNA sequence of CPn0349, which is the CP homologue of CT067 (ytgA), which was identified in clone CTL2gam18.

SEQ ID NO: 480 corresponds to the DNA sequence of CPn0325, which is the CP homologue of CT088 (sycE), which was identified in clone CTL2-19(5).

SEQ ID NO: 481 corresponds to the DNA sequence of CPn0326, which is the CP homologue of CT087 (malQ), which was identified in clone CTL2-19(5°).

SEQ ID NO: 482 corresponds to the DNA sequence of CPn0793, which is the CP homologue of CT588 (rbsu), which was identified in clone CTL2gam23.

SEQ ID NO: 483 corresponds to the DNA sequence of CPn0199, which is the CP homologue of CT199 (oppB1), which was identified in clone CTL2gam24.

SEQ ID NO: 484 corresponds to the DNA sequence of CPn0666, which is the CP homologue of CT545 (dnaE), which was identified in clone CTL2-24.

SEQ ID NO: 485 corresponds to the DNA sequence of CPn0065, which is the CP homologue of CT288, which was identified in clone CTL2gam27.

SEQ ID NO: 486 corresponds to the DNA sequence of CPn0444, which is the CP homologue of CT413 (pmpB), which was identified in clone CTL2gam30(5')(3').

SEQ ID NO: 487 corresponds to the DNA sequence of CPn-ORF5, which is the CP homologue of CT-ORF3, which was identified in clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5'), and CTL2-23.

SEQ ID NO: 488 corresponds to the DNA sequence of CPn-ORF6, which is the CP homologue of CT-ORF4, which was identified in clone CTL2-18(3').

SEQ ID NO: 489 corresponds to the DNA sequence of CP-ORF7, which is the CP homologue of CT-ORF5, which was identified in clone CTL2-18(3').

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SEQ ID NO: 490 corresponds to the amino acid sequence of CPn0894, which is the CP homologue of CT751 (amn), which was identified in clones CTL2-1 and CTL2-5.

SEQ ID NO: 491 corresponds to the amino acid sequence of CPn0074, which is the CP homologue of CT332 (tuf), which was identified in clone CTL2-2.

SEQ ID NO: 492 corresponds to the amino acid sequence of CPn0122, which is the CP homologue of CT032 (metG), which was identified in clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 493 corresponds to the amino acid sequence of CPn0121, which is the CP homologue of CT031, which was identified in clone CTL2-3(5')(3').

SEQ ID NO: 494 corresponds to the amino acid sequence of CPn0120 which is the CP homologue of CT030 (gmK) which was identified in clones CTL2-3 (3') and CTL2-21.

SEQ ID NO: 495 corresponds to the amino acid sequence of CPn0359, which is the CP homologue of CT064 (lepA), which was identified in clone CTL2gam5.

SEQ ID NO: 496 corresponds to the amino acid sequence of CPn0414, which is the CP homologue of CT265 (accA), which was identified in clone CTL2-6.

SEQ ID NO: 497 corresponds to the amino acid sequence of CPn0413, which is the CP homologue of CT264 (msbA), which was identified in clone CTL2-6.

SEQ ID NO: 498 corresponds to the amino acid sequence of CPn0394, which is the CP homologue of CT256, which was identified in clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 499 corresponds to the amino acid sequence of CPn0395, which is the CP homologue of CT257, which was identified in clones CTL2cam6(5') and CTL2-11(5').

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SEQ ID NO: 500 corresponds to the amino acid sequence of CPn0487, which is the CP homologue of CT384, which was identified in clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 501 corresponds to the amino acid sequence of CPn0592, which is the CP homologue of CT473, which was identified in clone CTL2-8b.

SEQ ID NO: 502 corresponds to the amino acid sequence of CPn0593, which is the CP homologue of CT474, which was identified in clone CTL2-8b.

SEQ ID NO: 503 corresponds to the amino acid sequence of CPn0197, which is the CP homologue of CT139 (oppA1), which was identified in clone CTL2-8b.

SEQ ID NO: 504 corresponds to the amino acid sequence of CPn0363, which is the CP homologue of CT060 (flhA), which was identified in clone CT1.2-8b.

SEQ ID NO: 505 corresponds to the amino acid sequence of CPn0301, which is the CP homologue of CT242, which was identified in clone CTL2gam8.

SEQ ID NO: 506 corresponds to the amino acid sequence of CPn0302, which is the CP homologue of CT243 (lpxD), which was identified in clone CTL2gam8.

SEQ ID NO: 507 corresponds to the amino acid sequence of CPn0324, which is the CP homologue of CT089 (IcrE), which was identified in clones CTL2-9, CTL2gam1, CTL2gam17 and CTL2-19(5*).

SEQ ID NO: 508 corresponds to the amino acid sequence of CPn0761, which is the CP homologue of CT610, which was identified in clone CTL2-10(5°)(3°).

SEQ ID NO: 509 corresponds to the amino acid sequence of CPn0760, which is the CP homologue of CT611, which was identified in clone CTL2-10(5*).

SEQ ID NO: 510 corresponds to the amino acid sequence of CPn0329, which is the CP homologue of CT154, which was identified in clones CTL2gam10 and CTL2gam21.

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SEQ ID NO: 511 corresponds to the amino acid sequence of CPn0990, which is the CP homologue of CT833 (infC), which was identified in clone CTL2-12.

SEQ ID NO: 512 corresponds to the amino acid sequence of CPn-ORF5, which is the CP homologue of CT ORF3, which was identified in clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5'), and CTL2-23.

SEQ ID NO: 513 corresponds to the amino acid sequence of CPn0984, which is the CP homologue of CT827 (nrdA) which was identified in clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 514 corresponds to the amino acid sequence of CPn0985, which is the CP homologue of CT828 (nrdB) which was identified in clones CTL2-16(3') CTL2gam15(3').

SEQ ID NO: 515 corresponds to the amino acid sequence of CPn0349, which is the CP homologue of CT067 (ytgA), which was identified in clone CTL2gam18.

SEQ ID NO: 516 corresponds to the DNA sequence of CPn-ORF6, which is the CP homologue of CT-ORF4, which was identified in clone CTL2-18(3').

SEQ ID NO: 517 corresponds to the DNA sequence of CP-ORF7, which is the CP homologue of CT-ORF5, which was identified in clone CTL2-18(3').

SEQ ID NO: 518 corresponds to the amino acid sequence of CPn0326, which is the CP homologue of CT087 (malQ), which was identified in clone CTL2-19(5').

SEQ ID NO: 519 corresponds to the amino acid sequence of CPn0325, which is the CP homologue of CT088 (sycE), which was identified in clone CTL2-19(5').

SEQ ID NO: 520 corresponds to the amino acid sequence of CPn0793, which is the CP homologue of CT588 (rbsu), which was identified in clone CTL2gam23.

SEQ ID NO: 521 corresponds to the amino acid sequence of CPn0199, which is the CP homologue of CT199 (oppB1), which was identified in clone CTL2gam24.

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SEQ ID NO: 522 corresponds to the amino acid sequence of CPn0666, which is the CP homologue of CT545 (dnaE), which was identified in clone CTL2-24.

SEQ ID NO: 523 corresponds to the DNA sequence of CPn0065, which is the CP homologue of CT288, which was identified in clone CTL2gam27.

SEQ ID NO: 524 corresponds to the DNA sequence of CPn0444, which is the CP homologue of CT413 (pmpB), which was identified in clone CTL2gam30(5'Y(3').

SEQ ID NO: 525 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT751 (amn) identified from the clones CTL2-1 and CTL2-5.

SEQ ID NO: 526 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT322 (tuff) identified from the clone CTL2-2.

SEQ ID NO: 527 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT032 (metG) identified from the clones CTL2gam2, CTL2-3(5') and CTL2-4.

SEQ ID NO: 528 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT031 identified from the clone CTL2-3(5')(3').

SEQ ID NO: 529 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT030 (gmK) identified from the clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 530 sets forth the full-length *C. trachomalis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT064 (lepA) identified from the clone CTL2gam5.

SEQ ID NO: 531 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT265 (accA) identified from the clone CTL2-6.

SEQ ID NO: 532 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT624 (msbA) identified from the clones CTL2-6.

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SEQ ID NO: 533 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT256 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 534 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT257 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 535 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT384 identified from the clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 536 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT473 identified from the clone CTL2-8b.

SEQ ID NO: 537 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT474 identified from the clones CTL2-8b.

SEQ ID NO: 538 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT139 (oppA1) identified from the clones CTL2-8b.

SEQ ID NO: 539 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT060 (flhA) identified from the clone CT1.2-8b.

SEQ ID NO: 540 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT242 identified from the clone CTL2gam8.

SEQ ID NO: 541 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT243 (lpxD) identified from the clone CTL2gam8.

SEQ ID NO: 542 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT089 identified from the clones CTL2-9, CTL2gam1, CTL2gam17, and CTL2-19(5').

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SEQ ID NO: 543 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT610 identified from the clone CTL2-10 (5°Y3°).

SEQ ID NO: 544 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT611 identified from the clone CTL2-10(5*).

SEQ ID NO: 545 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT154 identified from the clones CTL2gam10 and CTL2gam21.

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SEQ ID NO: 546 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT833 (infC) identified from the clone CTL2-12.

SEQ ID NO: 547 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT827 (nrdA) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 548 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT828 (nrdB) identified from the clones CTL2-16(3') and CTL2gam15(3').

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SEQ ID NO: 549 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT067 (ytgA) identified from the clone CTL2gam18.

SEQ ID NO: 550 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT088 (svcE) identified from the clones CTL2-19(5').

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SEQ ID NO: 551 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT087 identified from the clone CTL2-19(5').

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SEQ ID NO: 552 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT588 (rsbu) identified from the clone CTL2gam23.

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SEQ ID NO: 553 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT199 (oppB1) identified from the clone CTL2gam24.

SEQ ID NO: 554 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT545 (dnaE) identified from the clone CTL2-4.

SEQ ID NO: 555 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT288 identified from the clones CTL20am27.

SEQ ID NO: 556 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT413 (pmpB) identified from the clone CTL2gam30(5°)(3°).

SEQ ID NO: 557 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF3 identified from the clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5') and CTL2-23.

SEQ ID NO: 558 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for pCT-ORF4 identified from the clone CTL2-18(3').

SEQ ID NO: 559 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF5 identified from the clones CTL2-18(3').

SEQ ID NO: 560 sets forth the full-length *C. trachomatis* scrovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT751 (amn) identified from the clones CTL2-1 and CTL2-5.

SEQ ID NO: 561 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT322 (tuff) identified from the clone CTL2-2.

SEQ ID NO: 562 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT032 (metG) identified from the clones CTL2gam2, CTL2-3(5') and CTL2-4.

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SEQ ID NO: 563 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT031 identified from the clone CTL2-3(5')(3').

SEQ ID NO: 564 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT030 (gmK) identified from the clones CTL2-3(3') and CTL2-21.

SEQ ID NO: 565 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT064 (lepA) identified from the clone CTL2gam5.

SEQ ID NO: 566 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT265 (accA) identified from the clone CTL2-6.

SEQ ID NO: 567 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT624 (msbA) identified from the clones CTL2-6.

SEQ ID NO: 568 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT256 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 569 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT257 identified from the clones CTL2gam6(5') and CTL2-11(5').

SEQ ID NO: 570 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT384 identified from the clones CTL2gam6(3') and CTL2-11(3').

SEQ ID NO: 571 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT473 identified from the clone CTL2-8b.

SEQ ID NO: 572 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT474 identified from the clones CTL2-8b.

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SEQ ID NO: 573 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT139 (oppA1) identified from the clones CTL2-8b.

SEQ ID NO: 574 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT060 (flhA) identified from the clone CTL2-8b.

SEQ ID NO: 575 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT242 identified from the clone CTL2gam8.

SEQ ID NO: 576 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT243 (lpxD) identified from the clone CTL2gam8.

SEQ ID NO: 577 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT089 identified from the clones CTL2-9, CTL2gam1, CTL2gam17, and CTL2-19(5*).

SEQ ID NO: 578 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT610 identified from the clone CTL2-10 (5')(3').

SEQ ID NO: 579 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT611 identified from the clone CTL2-10(5').

SEQ ID NO: 580 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT154 identified from the clones CTL2gam10 and CTL2gam21.

SEQ ID NO: 581 sets forth the full-length *C. trachomatis* serovar D amino acid sequence homologous to the *C. trachomatis* LGV II sequence for CT833 (infC) identified from the clone CTL2-12.

SEQ ID NO: 582 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT-ORF3 identified from the clones CTL2gam15(5'), CTL2-16(5'), CTL2-18(5') and CTL2-23.

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SEO ID NO: 583 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT827 (nrdA) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEO ID NO: 584 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT828 (nrdB) identified from the clones CTL2-16(3') and CTL2gam15(3').

SEQ ID NO: 585 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT067 (vtgA) identified from the clone CTL2gam18.

SEQ ID NO: 586 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for pCT-ORF4 identified from the clone CTL2-18(3')

SEO ID NO: 587 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT-ORF5 identified from the clones CTL2-18(3').

SEQ ID NO: 588 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT087 identified from the clone CTL2-19(5').

SEO ID NO: 589 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT088 (sycE) identified from the clones CTL2-19(5').

SEO ID NO: 590 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT588 (rsbu) identified from the clone CTL2gam23.

SEO ID NO: 591 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT199 (oppB1) identified from the clone CTL2gam24.

SEQ ID NO: 592 sets forth the full-length C. trachomatis serovar D DNA sequence homologous to the C. trachomatis LGV II sequence for CT545 (dnaE) identified from the clone CTL2-4.

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SEQ ID NO: 593 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT288 identified from the clones CTL2gam27.

SEQ ID NO: 594 sets forth the full-length *C. trachomatis* serovar D DNA sequence homologous to the *C. trachomatis* LGV II sequence for CT413 (pmpB) identified from the clone CTL2gam30(5')(3').

SEQ ID NO: 595 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0406, of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 596 sets forth the DNA sequence for the Chlamydia pneumoniae homologue, CPn0315, of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 597 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0406, of the Chlamydia trachomatis gene CT102.

SEQ ID NO: 598 sets forth the amino acid sequence for the Chlamydia pneumoniae homologue, CPn0315, of the Chlamydia trachomatis gene CT098.

SEQ ID NO: 599 sets forth the amino acid sequence for Chlamydia trachomatis serovar D CT287 protein.

DESCRIPTION OF THE FIGURES

- Fig. 1 illustrates induction of INF-γ from a Chlamydia-specific T cell line activated by target cells expressing clone 4C9-18#2.
 - Fig. 2 illustrates retroviral vectors pBIB-KS1,2,3 modified to contain a Kosak translation initiation site and stop codons.
 - Fig. 3 shows specific lysis in a chromium release assay of P815 cells pulsed with Chlamydia peptides CtC7.8-12 (SEQ ID NO: 18) and CtC7.8-13 (SEQ ID NO: 19).
- 25 Fig. 4 shows antibody isotype titers in C57Bl/6 mice immunized with C. trachomatis SWIB protein.
 - Fig. 5 shows *Chlamydia*-specific T-cell proliferative responses in splenocytes from C3H mice immunized with *C. trachomatis* SWIB protein.
- Fig. 6 illustrates the 5' and 3' primer sequences designed from C. pneumoniae which
 were used to isolate the SWIB and S13 genes from C. pneumoniae.

Figs. 7A and 7B show induction of IFN-γ from a human anti-chlamydia T-cell line (TCL-8) capable of cross-reacting to C. trachomatis and C. pneumonia upon activation by monocyte-derived dendritic cells expressing chlamydial proteins.

Fig. 8 shows the identification of T cell epitopes in Chlamydial ribosomal S13 protein
with T-cell line TCL 8 EB/DC.

Fig. 9A and B illustrate the proliferative response of CP-21 T-cells generated against *C. pnuemoniae*-infected dendritic cells to recombinant *C. pneumonia-SWIB* protein, but not *C. trachomatis* SWIB protein.

Fig. 10 shows the *C. trachomatis*-specific SWIB proliferative responses of a primary T-10 cell line (TCT-10 EB) from an asymptomatic donor.

Fig. 11 illustrates the identification of T-cell epitope in *C. trachomatis* SWIB with an antigen specific T-cell line (TCL-10 EB).

Fig. 12 shows the C. trachomatis-specific proliferative responses of primary T cell lines generated from two patients against the CT specific antigens CT622, CT875 and CT 15 FB.

DETAILED DESCRIPTION OF THE INVENTION

As noted above, the present invention is generally directed to compositions and methods for the diagnosis and treatment of Chlamydial infection. In one aspect, the compositions of the subject invention include polypeptides that comprise at least one immunogenic portion of a Chlamydia antigen, or a variant thereof.

In specific embodiments, the subject invention discloses polypeptides comprising an immunogenic portion of a *Chlamydia* antigen, wherein the *Chlamydia* antigen comprises an amino acid sequence encoded by a polynucleotide molecule disclosed herein, the complements of said nucleotide sequences, and variants of such sequences.

As used herein, the term "polypeptide" encompasses amino acid chains of any length, including full length proteins (i.e., antigens), wherein the amino acid residues are linked by covalent peptide bonds. Thus, a polypeptide comprising an immunogenic portion of one of the inventive antigens may consist entirely of the immunogenic portion, or may contain additional sequences. The additional sequences

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may be derived from the native *Chlamydia* antigen or may be heterologous, and such sequences may (but need not) be immunogenic.

The term "polynucleotide(s)," as used herein, means a single or doublestranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and
corresponding RNA molecules, including HnRNA and mRNA molecules, both sense
and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA,
as well as wholly or partially synthesized polynucleotides. An HnRNA molecule
contains introns and corresponds to a DNA molecule in a generally one-to-one manner.
An mRNA molecule corresponds to an HnRNA and DNA molecule from which the
introns have been excised. A polynucleotide may consist of an entire gene, or any
portion thereof. Operable anti-sense polynucleotides may comprise a fragment of the
corresponding polynucleotide, and the definition of "polynucleotide" therefore includes
all such operable anti-sense fragments.

An "immunogenic portion" of an antigen is a portion that is capable of reacting with sera obtained from a Chlamydia-infected individual (i.e., generates an absorbance reading with sera from infected individuals that is at least three standard deviations above the absorbance obtained with sera from uninfected individuals, in a representative ELISA assay described herein). Such immunogenic portions generally comprise at least about 5 amino acid residues, more preferably at least about 10, and 20 most preferably at least about 20 amino acid residues. Methods for preparing and identifying immunogenic portions of antigens of known sequence are well known in the art and include those summarized in Paul, Fundamental Immunology, 3rd ed., Raven Press, 1993, pp. 243-247 and references cited therein. Such techniques include screening polypeptides for the ability to react with antigen-specific antibodies, antisera 25 and/or T-cell lines or clones. As used herein, antisera and antibodies are "antigenspecific" if they specifically bind to an antigen (i.e., they react with the protein in an ELISA or other immunoassay, and do not react detectably with unrelated proteins). Such antisera and antibodies may be prepared as described herein, and using well known techniques. An immunogenic portion of a native Chlamydia protein is a portion 30 that reacts with such antisera and/or T-cells at a level that is not substantially less than the reactivity of the full length polypeptide (e.g., in an ELISA and/or T-cell reactivity assay). Such immunogenic portions may react within such assays at a level that is

similar to or greater than the reactivity of the full length polypeptide. Such screens may generally be performed using methods well known to those of ordinary skill in the art, such as those described in Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988. For example, a polypeptide may be immobilized on a solid support and contacted with patient sera to allow binding of antibodies within the sera to the immobilized polypeptide. Unbound sera may then be removed and bound antibodies detected using, for example, ¹²⁵I-labeled Protein A.

Examples of immunogenic portions of antigens contemplated by the present invention include, for example, the T cell stimulating epitopes provided in SEQ ID NO: 9, 10, 18, 19, 31, 39, 93-96, 98, 100-102, 106, 108, 138-140, 158, 167, 168, 246, 247 and 254-256. Polypeptides comprising at least an immunogenic portion of one or more *Chlamydia* antigens as described herein may generally be used, alone or in combination, to detect Chlamydial infection in a patient.

The compositions and methods of the present invention also encompass

variants of the above polypeptides and polynucleotide molecules. Such variants
include, but are not limited to, naturally occurring allelic variants of the inventive
scquences. In particular, variants include other *Chlamydiae* serovars, such as serovars

D, E and F, as well as the several LGV serovars which share homology to the inventive
polypeptide and polynucleotide molecules described herein. Preferably, the serovar

homologues show 95-99% homology to the corresponding polypeptide sequence(s)
described herein.

A polypeptide "variant," as used herein, is a polypeptide that differs from the recited polypeptide only in conservative substitutions and/or modifications, such that the antigenic properties of the polypeptide are retained. In a preferred embodiment, variant polypeptides differ from an identified sequence by substitution, deletion or addition of five amino acids or fewer. Such variants may generally be identified by modifying one of the above polypeptide sequences, and evaluating the antigenic properties of the modified polypeptide using, for example, the representative procedures described herein. In other words, the ability of a variant to react with antigen-specific antisera may be enhanced or unchanged, relative to the native protein, or may be diminished by less than 50%, and preferably less than 20%, relative to the native protein. Such variants may generally be identified by modifying one of the above

polypeptide sequences and evaluating the reactivity of the modified polypeptide with antigen-specific antibodies or antisera as described herein. Preferred variants include those in which one or more portions, such as an N-terminal leader sequence or transmembrane domain, have been removed. Other preferred variants include variants in which a small portion (e.g., 1-30 amino acids, preferably 5-15 amino acids) has been removed from the N- and/or C-terminal of the mature protein.

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As used herein, a "conservative substitution" is one in which an amino acid is substituted for another amino acid that has similar properties, such that one skilled in the art of peptide chemistry would expect the secondary structure and hydropathic nature of the polypeptide to be substantially unchanged. Amino acid substitutions may generally be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity and/or the amphipathic nature of the residues. For example, negatively charged amino acids include aspartic acid and glutamic acid; positively charged amino acids include lysine and arginine; and amino acids with uncharged polar head groups having similar hydrophilicity values include leucine, isoleucine and valine; glycine and alanine; asparagine and glutamine; and serine, threonine, phenylalanine and tyrosine. Other groups of amino acids that may represent conservative changes include: (1) ala, pro, gly, glu, asp, gln, asn, ser, thr; (2) cys, ser, tyr, thr; (3) val, ile, leu, met, ala, phe; (4) lys, arg, his; and (5) phe, tyr, trp, his. A variant may also, or alternatively, contain nonconservative changes. In a preferred embodiment, variant polypeptides differ from a native sequence by substitution, deletion or addition of five amino acids or fewer. Variants may also (or alternatively) be modified by, for example, the deletion or addition of amino acids that have minimal influence on the immunogenicity, secondary structure and hydropathic nature of the polypeptide. Variants may also, or alternatively, contain other modifications, including the deletion or addition of amino acids that have minimal influence on the antigenic properties, secondary structure and hydropathic nature of the polypeptide. For example, a polypeptide may be conjugated to a signal (or leader) sequence at the N-terminal end of the protein which co-translationally or post-translationally directs transfer of the protein. The polypeptide may also be conjugated to a linker or other sequence for ease of synthesis, purification or identification of the polypeptide (e.g., poly-His), or to WO 02/08267 PCT/US01/23121

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enhance binding of the polypeptide to a solid support. For example, a polypeptide may be conjugated to an immunoglobulin Fc region.

A polynucleotide "variant" is a sequence that differs from the recited nucleotide sequence in having one or more nucleotide deletions, substitutions or 5 additions such that the immunogenicity of the encoded polypeptide is not diminished, relative to the native protein. The effect on the immunogenicity of the encoded polypeptide may generally be assessed as described herein. Such modifications may be readily introduced using standard mutagenesis techniques, such as oligonucleotidedirected site-specific mutagenesis as taught, for example, by Adelman et al. (DNA, 2:183, 1983). Nucleotide variants may be naturally occurring allelic variants as discussed below, or non-naturally occurring variants. The polypeptides provided by the present invention include variants that are encoded by polynucleotide sequences which are substantially homologous to one or more of the polynucleotide sequences specifically recited herein. "Substantial homology," as used herein, refers to polynucleotide sequences that are capable of hybridizing under moderately stringent conditions. Suitable moderately stringent conditions include prewashing in a solution of 5X SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0); hybridizing at 50°C-65°C, 5X SSC, overnight or, in the event of cross-species homology, at 45°C with 0.5X SSC; followed by washing twice at 65°C for 20 minutes with each of 2X, 0.5X and 0.2X SSC containing 0.1% SDS. Such hybridizing polynucleotide sequences are also within the scope of this invention, as are nucleotide sequences that, due to code degeneracy, encode a polypeptide that is the same as a polypeptide of the present invention.

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Two nucleotide or polypeptide sequences are said to be "identical" if the sequence of nucleotides or amino acid residues in the two sequences is the same when aligned for maximum correspondence as described below. Comparisons between two sequences are typically performed by comparing the sequences over a comparison window to identify and compare local regions of sequence similarity. A "comparison window" as used herein, refers to a segment of at least about 20 contiguous positions, usually 30 to about 75, 40 to about 50, in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned.

Optimal alignment of sequences for comparison may be conducted using the Megalign program in the Lasergene suite of bioinformatics software (DNASTAR, Inc., Madison, WI), using default parameters. This program embodies several alignment schemes described in the following references: Dayhoff, M.O. (1978) A model of evolutionary change in proteins - Matrices for detecting distant relationships. In Dayhoff, M.O. (ed.) Atlas of Protein Sequence and Structure, National Biomedical Resarch Foundaiton, Washington DC Vol. 5, Suppl. 3, pp. 345-358; Hein J. (1990) Unified Approach to Alignment and Phylogenes pp. 626-645 Methods in Enzymology vol. 183, Academic Press, Inc., San Diego, CA; Higgins, D.G. and Sharp, P.M. (1989) Fast and sensitive multiple sequence alignments on a microcomputer CABIOS 5:151-153; Myers, E.W. and Muller W. (1988) Optimal alignments in linear space CABIOS 4:11-17; Robinson, E.D. (1971) Comb. Theor 11:105; Santou, N. Nes, M. (1987) The neighbor joining method. A new method for reconstructing phylogenetic trees Mol. Biol. Evol. 4:406-425; Sneath, P.H.A. and Sokal, R.R. (1973) Numerical Taxonomy -15 the Principles and Practice of Numerical Taxonomy, Freeman Press, San Francisco. CA; Wilbur, W.J. and Lipman, D.J. (1983) Rapid similarity searches of nucleic acid and protein data banks Proc. Natl. Acad., Sci. USA 80:726-730.

Alternatively, optimal alignment of sequences for comparison may be conducted by the local identity algorithm of Smith and Waterman (1981) Add. APL.

Math 2:482, by the identity alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity methods of Pearson and Lipman (1988) Proc. Natl. Acad. Sci. (U.S.A.) 85: 2444, by computerized implementations of these algorithms (GAP, BESTFIT, BLAST, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group (GCG), 575 Science Dr., Madison, WI), or by inspection.

One illustrative example of algorithms that are suitable for determining percent sequence identity and sequence similarity are the BLAST and BLAST 2.0 algorithms, which are described in Altschul et al. (1977) Nuc. Acids Res. 25:3389-3402 and Altschul et al. (1990) J. Mol. Biol. 215:403-410, respectively. BLAST and BLAST 2.0 can be used, for example with the parameters described herein, to determine percent sequence identity for the polynucleotides and polypeptides of the invention. Software for performing BLAST analyses is publicly available through the National Center for

Biotechnology Information (http://www.ncbi.nlm.nih.gov/) In one illustrative example, cumulative scores can be calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always >0) and N (penalty score for mismatching residues; always <0). For amino acid sequences, a scoring matrix can be used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, and expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915) alignments, (B) of 50, expectation (E) of 10, M=5, N=-4 and a comparison of both strands.

Preferably, the "percentage of sequence identity" is determined by comparing two optimally aligned sequences over a window of comparison of at least 20 positions, wherein the portion of the polynucleotide or amino acid sequence in the comparison window may comprise additions or deletions (i.e. gaps) of 20 percent or less, usually 5 to 15 percent, or 10 to 12 percent, as compared to the reference sequences (which does not comprise additions or deletions) for optimal alignment of the two sequences. The percentage is calculated by determining the number of positions at which the identical nucleic acid bases or amino acid residue occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the reference sequence (i.e. the window size) and multiplying the results by 100 to yield the percentage of sequence identity.

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Therefore, the present invention provides polynucleotide and polypeptide sequences having substantial identity to the sequences disclosed herein, for example those comprising at least 50% or more sequence identity, preferably at least 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% or higher, sequence identity compared to a polynucleotide or polypeptide sequence of this invention using the methods described herein, (e.g., BLAST analysis using standard parameters, as described below). One skilled in this art will recognize that these values can be

appropriately adjusted to determine corresponding identity of proteins encoded by two polynucleotide sequences by taking into account codon degeneracy, amino acid similarity, reading frame positioning and the like.

In additional embodiments, the present invention provides isolated 5 polynucleotides or polypeptides comprising various lengths of contiguous stretches of sequence identical to or complementary to one or more of the sequences disclosed herein. For example, polynucleotides and polyneptides encompassed by this invention may comprise at least about 15, 20, 30, 40, 50, 75, 100, 150, 200, 300, 400, 500 or 1000 or more contiguous nucleotides of one or more of the disclosed sequences, as well as all 10 intermediate lengths therebetween. It will be readily understood that "intermediate lengths", in this context, means any length between the quoted values, such as 16, 17, 18, 19, etc.; 21, 22, 23, etc.; 30, 31, 32, etc.; 50, 51, 52, 53, etc.; 100, 101, 102, 103, etc.; 150, 151, 152, 153, etc.; including all integers through the 200-500; 500-1.000, and the like.

The polynucleotides of the present invention, or fragments thereof, regardless of the length of the coding sequence itself, may be combined with other DNA sequences, such as promoters, polyadenylation signals, additional restriction enzyme sites, multiple cloning sites, other coding segments, and the like, such that their overall length may vary considerably. It is therefore contemplated that a nucleic acid fragment 20 of almost any length may be employed, with the total length preferably being limited by the ease of preparation and use in the intended recombinant DNA protocol. For example, illustrative DNA segments with total lengths of about 10,000, about 5000, about 3000, about 2,000, about 1,000, about 500, about 200, about 100, about 50 base pairs in length, and the like, (including all intermediate lengths) are contemplated to be 25 useful in many implementations of this invention.

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Also included in the scope of the present invention are alleles of the genes encoding the nucleotide sequences recited in herein. As used herein, an "allele" or "allellic sequence" is an alternative form of the gene which may result from at least one mutation in the nucleic acid sequence. Alleles may result in altered mRNAs or 30 polypeptides whose structure or function may or may not be altered. Any given gene may have none, one, or many allelic forms. Common mutational changes which give rise to alleles are generally ascribed to natural deletions, additions, or substitutions of nucleotides. Each of these types of changes may occur alone or in combination with the others, one or more times in a given sequence.

In specific embodiments, the subject invention discloses polypeptides comprising at least an immunogenic portion of a Chlamydia antigen (or a variant of 5 such an antigen), that comprises one or more of the amino acid sequences encoded by (a) a polynucleotide sequence selected from the group consisting of SEQ ID NO: 358-361, 407-430, 525-559, 582-598; (b) the complements of such DNA sequences or (c) DNA sequences substantially homologous to a sequence in (a) or (b). As discussed in the Examples below, several of the Chlamydia antigens disclosed herein recognize a 10 T cell line that recognizes both Chlamydia trachomatis and Chlamydia pneumoniae infected monocyte-derived dendritic cells, indicating that they may represent an immunoreactive epitope shared by Chlamvdia trachomatis and Chlamvdia pneumoniae. The antigens may thus be employed in a vaccine for both C. trachomatis genital tract infections and for C. pneumonia infections. Further characterization of these Chlamydia 15 antigens from Chlamvdia trachomatis and Chlamvdia pneumonia to determine the extent of cross-reactivity is provided in Example 6. Additionally, Example 4 describes cDNA fragments (SEQ ID NO: 15, 16 and 33) isolated from C. trachomatis which encode proteins (SEQ ID NO: 17-19 and 32) capable of stimulating a Chlamydiaspecific murine CD8+ T cell line.

In general, Chlamydia antigens, and polynucleotide sequences encoding such antigens, may be prepared using any of a variety of procedures. For example, polynucleotide molecules encoding Chlamydia antigens may be isolated from a Chlamvdia genomic or cDNA expression library by screening with a Chlamvdiaspecific T cell line as described below, and sequenced using techniques well known to 25 those of skill in the art. Additionally, a polynucleotide may be identified, as described in more detail below, by screening a microarray of cDNAs for Chlamydia-associated expression (i.e., expression that is at least two fold greater in Chlamydia-infected cells than in controls, as determined using a representative assay provided herein). Such screens may be performed using a Synteni microarray (Palo Alto, CA) according to the 30 manufacturer's instructions (and essentially as described by Schena et al., Proc. Natl. Acad. Sci. USA 93:10614-10619, 1996 and Heller et al., Proc. Natl. Acad. Sci. USA 94:2150-2155, 1997). Alternatively, polypeptides may be amplified from cDNA

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prepared from cells expressing the proteins described herein. Such polynucleotides may be amplified via polymerase chain reaction (PCR). For this approach, sequence-specific primers may be designed based on the sequences provided herein, and may be purchased or synthesized.

Antigens may be produced recombinantly, as described below, by inserting a polynucleotide sequence that encodes the antigen into an expression vector and expressing the antigen in an appropriate host. Antigens may be evaluated for a desired property, such as the ability to react with sera obtained from a Chlamydia-infected individual as described herein, and may be sequenced using, for example, traditional Edman chemistry. See Edman and Berg. Eur. J. Biochem. 80:116-132, 1967.

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Polynucleotide sequences encoding antigens may also be obtained by screening an appropriate Chlamydia cDNA or genomic DNA library for polynucleotide sequences that hybridize to degenerate oligonucleotides derived from partial amino acid sequences of isolated antigens. Degenerate oligonucleotide sequences for use in such a screen may be designed and synthesized, and the screen may be performed, as described (for example) in Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY (and references cited therein). Polymerase chain reaction (PCR) may also be employed, using the above oligonucleotides in methods well known in the art, to isolate a nucleic acid probe from a cDNA or genomic library. The library screen may then be performed using the isolated probe

An amplified portion may be used to isolate a full length gene from a suitable library (e.g., a Chlamydia cDNA library) using well known techniques. Within such techniques, a library (cDNA or genomic) is screened using one or more polynucleotide probes or primers suitable for amplification. Preferably, a library is size-selected to include larger molecules. Random primed libraries may also be preferred for identifying 5' and upstream regions of genes. Genomic libraries are preferred for obtaining introns and extending 5' sequences.

For hybridization techniques, a partial sequence may be labeled (e.g., by nick-translation or end-labeling with ³²P) using well known techniques. A bacterial or bacteriophage library is then screened by hybridizing filters containing denatured bacterial colonies (or lawns containing phage plaques) with the labeled probe (see

Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY, 1989). Hybridizing colonies or plaques are selected and expanded, and the DNA is isolated for further analysis. cDNA clones may be analyzed to determine the amount of additional sequence by, for example, PCR using a primer from the partial sequence and a primer from the vector. Restriction maps and partial sequences may be generated to identify one or more overlapping clones. The complete sequence may then be determined using standard techniques, which may involve generating a series of deletion clones. The resulting overlapping sequences are then assembled into a single contiguous sequence. A full length cDNA molecule can be generated by ligating suitable fragments, using well known techniques.

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Alternatively, there are numerous amplification techniques for obtaining a full length coding sequence from a partial cDNA sequence. Within such techniques, amplification is generally performed via PCR. Any of a variety of commercially available kits may be used to perform the amplification step. Primers may be designed using techniques well known in the art (see, for example, Mullis et al., Cold Spring Harbor Symp. Quant. Biol. 51:263, 1987; Erlich ed., PCR Technology, Stockton Press, NY, 1989), and software well known in the art may also be employed. Primers are preferably 22-30 nucleotides in length, have a GC content of at least 50% and anneal to the target sequence at temperatures of about 68°C to 72°C. The amplified region may be sequenced as described above, and overlapping sequences assembled into a continuous sequence.

One such amplification technique is inverse PCR (see Triglia et al., Nucl. Acids Res. 16:8186, 1988), which uses restriction enzymes to generate a fragment in the known region of the gene. The fragment is then circularized by intramolecular ligation and used as a template for PCR with divergent primers derived from the known region. Within an alternative approach, sequences adjacent to a partial sequence may be retrieved by amplification with a primer to a linker sequence and a primer specific to a known region. The amplified sequences are typically subjected to a second round of amplification with the same linker primer and a second primer specific to the known region. A variation on this procedure, which employs two primers that initiate extension in opposite directions from the known sequence, is described in WO 96/38591. Additional techniques include capture PCR (Lagerstrom et al., PCR Methods

Applic, 1:111-19, 1991) and walking PCR (Parker et al., Nucl. Acids. Res. 19:3055-60, 1991). Transcription-Mediated Amplification, or TMA is another method that may be utilized for the amplification of DNA, rRNA, or mRNA, as described in Patent No. PCT/US91/03184. This autocatalytic and isothermic non-PCR based method utilizes 5 two primers and two enzymes: RNA polymerase and reverse transcriptase. One primer contains a promoter sequence for RNA polymerase. In the first amplification, the promoter-primer hybridizes to the target rRNA at a defined site. Reverse transcriptase creates a DNA copy of the target rRNA by extension from the 3'end of the promoterprimer. The RNA in the resulting complex is degraded and a second primer binds to the 10 DNA copy. A new strand of DNA is synthesized from the end of the primer by reverse transcriptase creating double stranded DNA. RNA polymerase recognizes the promoter sequence in the DNA template and initiates transcription. Each of the newly synthesized RNA amplicons re-enters the TMA process and serves as a template for a new round of replication leading to the expotential expansion of the RNA amplicon. Other methods employing amplification may also be employed to obtain a full length cDNA sequence.

In certain instances, it is possible to obtain a full length cDNA sequence by analysis of sequences provided in an expressed sequence tag (EST) database, such as that available from GenBank. Searches for overlapping ESTs may generally be performed using well known programs (e.g., NCBI BLAST searches), and such ESTs may be used to generate a contiguous full length sequence. Full length cDNA sequences may also be obtained by analysis of genomic fragments.

Polynucleotide variants may generally be prepared by any method known in the art, including chemical synthesis by, for example, solid phase phosphoramidite chemical synthesis. Modifications in a polynucleotide sequence may also be introduced using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis (see Adelman et al., DNA 2:183, 1983). Alternatively, RNA molecules may be generated by in vitro or in vivo transcription of DNA sequences encoding a Chlamydial protein, or portion thereof, provided that the DNA is incorporated into a vector with a suitable RNA polymerase promoter (such as T7 or SP6). Certain portions may be used to prepare an encoded polypeptide, as described herein. In addition, or alternatively, a portion may be administered to a patient such that the encoded

polypeptide is generated in vivo (e.g., by transfecting antigen-presenting cells, such as dendritic cells, with a cDNA construct encoding a Chlamydial polypeptide, and administering the transfected cells to the patient).

A portion of a sequence complementary to a coding sequence (*i.e.*, an satisense polynucleotide) may also be used as a probe or to modulate gene expression. cDNA constructs that can be transcribed into antisense RNA may also be introduced into cells of tissues to facilitate the production of antisense RNA. An antisense polynucleotide may be used, as described herein, to inhibit expression of a *Chlamydial* protein. Antisense technology can be used to control gene expression through triple-helix formation, which compromises the ability of the double helix to open sufficiently for the binding of polymerases, transcription factors or regulatory molecules (*see* Gee et al., *In* Huber and Carr, *Molecular and Immunologic Approaches*, Futura Publishing Co. (Mt. Kisco, NY; 1994)). Alternatively, an antisense molecule may be designed to hybridize with a control region of a gene (*e.g.*, promoter, enhancer or transcription initiation site), and block transcription of the gene; or to block translation by inhibiting binding of a transcript to ribosomes.

A portion of a coding sequence, or of a complementary sequence, may also be designed as a probe or primer to detect gene expression. Probes may be labeled with a variety of reporter groups, such as radionuclides and enzymes, and are preferably at least 10 nucleotides in length, more preferably at least 20 nucleotides in length and still more preferably at least 30 nucleotides in length. Primers, as noted above, are preferably 22-30 nucleotides in length.

Any polynucleotide may be further modified to increase stability in vivo.

Possible modifications include, but are not limited to, the addition of flanking
sequences at the 5' and/or 3' ends; the use of phosphorothioate or 2' O-methyl rather
than phosphodiesterase linkages in the backbone; and/or the inclusion of nontraditional
bases such as inosine, queosine and wybutosine, as well as acetyl- methyl-, thio- and
other modified forms of adenine, cytidine, guanine, thymine and uridine.

Nucleotide sequences as described herein may be joined to a variety of
other nucleotide sequences using established recombinant DNA techniques. For
example, a polynucleotide may be cloned into any of a variety of cloning vectors,
including plasmids, phagemids, lambda phage derivatives and cosmids. Vectors of

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particular interest include expression vectors, replication vectors, probe generation vectors and sequencing vectors. In general, a vector will contain an origin of replication functional in at least one organism, convenient restriction endonuclease sites and one or more selectable markers. Other elements will depend upon the desired use, and will be apparent to those of ordinary skill in the art.

Synthetic polypeptides having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using techniques well known in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as the Merrifield solid-phase synthesis method, where amino acids are sequentially added to a growing amino acid chain. See Merrifield, J. Am. Chem. Soc. 85:2149-2146, 1963. Equipment for automated synthesis of polypeptides is commercially available from suppliers such as Perkin Elmer/Applied BioSystems Division, Foster City, CA, and may be operated according to the manufacturer's instructions.

As noted above, immunogenic portions of Chlamydia antigens may be prepared and identified using well known techniques, such as those summarized in Paul, Fundamental Immunology, 3d ed., Raven Press, 1993, pp. 243-247 and references cited therein. Such techniques include screening polypeptide portions of the native antigen for immunogenic properties. The representative ELISAs described herein may generally be employed in these screens. An immunogenic portion of a polypeptide is a portion that, within such representative assays, generates a signal in such assays that substantially similar to that generated by the full length antigen. In other words, an immunogenic portion of a Chlamydia antigen generates at least about 20%, and preferably about 100%, of the signal induced by the full length antigen in a model ELISA as described herein.

Portions and other variants of *Chlamydia* antigens may be generated by synthetic or recombinant means. Variants of a native antigen may generally be prepared using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis. Sections of the polynucleotide sequence may also be removed using standard techniques to permit preparation of truncated polyneptides.

Recombinant polypeptides containing portions and/or variants of a native antigen may be readily prepared from a polynucleotide sequence encoding the

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polypeptide using a variety of techniques well known to those of ordinary skill in the art. For example, supernatants from suitable host/vector systems which secrete recombinant protein into culture media may be first concentrated using a commercially available filter. Following concentration, the concentrate may be applied to a suitable purification matrix such as an affinity matrix or an ion exchange resin. Finally, one or more reverse phase HPLC steps can be employed to further purify a recombinant protein.

Any of a variety of expression vectors known to those of ordinary skill in the art may be employed to express recombinant polypeptides as described herein.

Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polynucleotide molecule that encodes a recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are E. coli, yeast or a mammalian cell line, such as COS or CHO. The DNA sequences expressed in this manner may encode naturally occurring antigens, portions of naturally occurring antigens, or other variants thereof.

In general, regardless of the method of preparation, the polypeptides disclosed herein are prepared in an isolated, substantially pure, form. Preferably, the polypeptides are at least about 80% pure, more preferably at least about 90% pure and most preferably at least about 99% pure.

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Within certain specific embodiments, a polypeptide may be a fusion protein that comprises multiple polypeptides as described herein, or that comprises at least one polypeptide as described herein and an unrelated sequence, such as a known Chlamydial protein. A fusion partner may, for example, assist in providing T helper epitopes (an immunological fusion partner), preferably T helper epitopes recognized by humans, or may assist in expressing the protein (an expression enhancer) at higher yields than the native recombinant protein. Certain preferred fusion partners are both immunological and expression enhancing fusion partners. Other fusion partners may be selected so as to increase the solubility of the protein or to enable the protein to be targeted to desired intracellular compartments. Still further fusion partners include affinity tags, which facilitate purification of the protein. A DNA sequence encoding a fusion protein of the present invention may be constructed using known recombinant

and the second polypeptides.

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DNA techniques to assemble separate DNA sequences encoding, for example, the first and second polypeptides, into an appropriate expression vector. The 3' end of a DNA sequence encoding the first polypeptide is ligated, with or without a peptide linker, to the 5' end of a DNA sequence encoding the second polypeptide so that the reading frames of the sequences are in phase to permit mRNA translation of the two DNA sequences into a single fusion protein that retains the biological activity of both the first

A peptide linker sequence may be employed to separate the first and the second polypeptides by a distance sufficient to ensure that each polypeptide folds into 10 its secondary and tertiary structures. Such a peptide linker sequence is incorporated into the fusion protein using standard techniques well known in the art. Suitable peptide linker sequences may be chosen based on the following factors: (1) their ability to adopt a flexible extended conformation; (2) their inability to adopt a secondary structure that could interact with functional epitopes on the first and second polypeptides; and 15 (3) the lack of hydrophobic or charged residues that might react with the polypeptide functional epitopes. Preferred peptide linker sequences contain Gly. Asn and Ser residues. Other near neutral amino acids, such as Thr and Ala may also be used in the linker sequence. Amino acid sequences which may be usefully employed as linkers include those disclosed in Maratea et al., Gene 40:39-46, 1985; Murphy et al., Proc. Natl. Acad. Sci. USA 83:8258-8562, 1986; U.S. Patent No. 4,935,233 and U.S. Patent No. 4.751,180. The linker sequence may be from 1 to about 50 amino acids in length. As an alternative to the use of a peptide linker sequence (when desired), one can utilize non-essential N-terminal amino acid regions (when present) on the first and second polypeptides to separate the functional domains and prevent steric hindrance.

The ligated DNA sequences are operably linked to suitable transcriptional or translational regulatory elements. The regulatory elements responsible for expression of DNA are located only 5' to the DNA sequence encoding the first polypeptides. Similarly, stop codons required to end translation and transcription termination signals are only present 3' to the DNA sequence encoding the second polypeptide.

Fusion proteins are also provided that comprise a polypeptide of the present invention together with an unrelated immunogenic protein. Preferably the WO 02/08267 PCT/US01/23121 64

immunogenic protein is capable of eliciting a recall response. Examples of such proteins include tetanus, tuberculosis and hepatitis proteins (see, for example, Stoute et al. New Engl. J. Med., 336:86-91, 1997).

Within preferred embodiments, an immunological fusion partner is 5 derived from protein D, a surface protein of the gram-negative bacterium Haemophilus influenza B (WO 91/18926). Preferably, a protein D derivative comprises approximately the first third of the protein (e.g., the first N-terminal 100-110 amino acids), and a protein D derivative may be lipidated. Within certain preferred embodiments, the first 109 residues of a Lipoprotein D fusion partner is included on the 10 N-terminus to provide the polypeptide with additional exogenous T-cell epitopes and to increase the expression level in E. coli (thus functioning as an expression enhancer). The lipid tail ensures optimal presentation of the antigen to antigen presenting cells. Other fusion partners include the non-structural protein from influenzae virus, NS1 (hemaglutinin). Typically, the N-terminal 81 amino acids are used, although different fragments that include T-helper epitopes may be used.

In another embodiment, the immunological fusion partner is the protein known as LYTA, or a portion thereof (preferably a C-terminal portion). LYTA is derived from Streptococcus pneumoniae, which synthesizes an N-acetyl-L-alanine amidase known as amidase LYTA (encoded by the LytA gene; Gene 43:265-292, 1986). LYTA is an autolysin that specifically degrades certain bonds in the peptidoglycan backbone. The C-terminal domain of the LYTA protein is responsible for the affinity to the choline or to some choline analogues such as DEAE. This property has been exploited for the development of E. coli C-LYTA expressing plasmids useful for expression of fusion proteins. Purification of hybrid proteins containing the C-LYTA fragment at the amino terminus has been described (see Biotechnology 10:795-798, 25 1992). Within a preferred embodiment, a repeat portion of LYTA may be incorporated into a fusion protein. A repeat portion is found in the C-terminal region starting at residue 178. A particularly preferred repeat portion incorporates residues 188-305.

In another embodiment, a Mycobacterium tuberculosis-derived Ra12 polynucleotide is linked to at least an immunogenic portion of a polynucleotide of this invention. Ra12 compositions and methods for their use inenhancing expression of heterologous polynucleotide sequences is described in U.S. Patent Application WO 02/08267 PCT/US01/23121

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60/158,585, the disclosure of which is incorporated herein by reference in its entirety. Briefly, Ra12 refers to a polynucleotide region that is a subsequence of a Mycobacterium tuberculosis MTB32A nucleic acid. MTB32A is a serine protease of 32 KD molecular weight encoded by a gene in virulent and avirulent strains of M. 5 tuberculosis. The nucleotide sequence and amino acid sequence of MTB32A have been described (U.S. Patent Application 60/158,585; see also, Skeiky et al., Infection and Immun. (1999) 67:3998-4007, incorporated herein by reference. In one embodiment, the Ra12 polypeptide used in the production of fusion polypeptides comprises a Cterminal fragment of the MTB32A coding sequence that is effective for enhancing the 10 expression and/or immunogenicity of heterologous Chlamydial antigenic polypeptides with which it is fused. In another embodiment, the Ra12 polypeptide corresponds to an approximately 14 kD. C-terminal fragment of MTB32A comprising some or all of amino acid residues 192 to 323 of MTB32A.

Recombinant nucleic acids, which encode a fusion polypeptide 15 comprising a Ra12 polypeptide and a heterologous Chlamydia polypeptide of interest, can be readily constructed by conventional genetic engineering techniques. Recombinant nucleic acids are constructed so that, preferably, a Ra12 polynucleotide sequence is located 5' to a selected heterologous Chlamydia polynucleotide sequence. It may also be appropriate to place a Ra12 polynucleotide sequence 3' to a selected heterologous polynucleotide sequence or to insert a heterologous polynucleotide sequence into a site within a Ra12 polynucleotide sequence.

In addition, any suitable polynucleotide that encodes a Ra12 or a portion or other variant thereof can be used in constructing recombinant fusion polynucleotides comprising Ra12 and one or more Chlamydia polynucleotides disclosed herein. Preferred Ra12 polynucleotides generally comprise at least about 15 consecutive nucleotides, at least about 30 nucleotides, at least about 60 nucleotides, at least about 100 nucleotides, at least about 200 nucleotides, or at least about 300 nucleotides that encode a portion of a Ra12 polypeptide.

Ra12 polynucleotides may comprise a native sequence (i.e., an endogenous sequence that encodes a Ra12 polypeptide or a portion thereof) or may comprise a variant of such a sequence. Ra12 polynucleotide variants may contain one WO 02/08267 PCT/US01/23121

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or more substitutions, additions, deletions and/or insertions such that the biological activity of the encoded fusion polypeptide is not substantially diminished, relative to a fusion polypeptide comprising a native Ra12 polypeptide. Variants preferably exhibit at least about 70% identity, more preferably at least about 80% identity and most preferably at least about 90% identity to a polypucleotide sequence that encodes a native Ra12 polypeptide or a portion thereof.

In another aspect, the present invention provides methods for using one or more of the above polypeptides or fusion proteins (or polynucleotides encoding such polypeptides or fusion proteins) to induce protective immunity against Chlamydial infection in a patient. As used herein, a "patient" refers to any warm-blooded animal, preferably a human. A patient may be afflicted with a disease, or may be free of detectable disease and/or infection. In other words, protective immunity may be induced to prevent or treat Chlamydial infection.

In this aspect, the polypeptide, fusion protein or polynucleotide molecule
is generally present within a pharmaceutical composition or a vaccine. Pharmaceutical
compositions may comprise one or more polypeptides, each of which may contain one
or more of the above sequences (or variants thereof), and a physiologically acceptable
carrier. Vaccines may comprise one or more of the above polypeptides and an
immunostimulant, such as an adjuvant or a liposome (into which the polypeptide is
incorporated). Such pharmaceutical compositions and vaccines may also contain other
Chlamydia antigens, either incorporated into a combination polypeptide or present
within a separate polypeptide.

Alternatively, a vaccine may contain polynucleotides encoding one or more polypeptides or fusion proteins as described above, such that the polypeptide is generated in situ. In such vaccines, the polynucleotides may be present within any of a variety of delivery systems known to those of ordinary skill in the art, including nucleic acid expression systems, bacterial and viral expression systems. Appropriate nucleic acid expression systems contain the necessary polynucleotide sequences for expression in the patient (such as a suitable promoter and terminating signal). Bacterial delivery systems involve the administration of a bacterium (such as Bacillus-Calmette-Guerrin) that expresses an immunogenic portion of the polypentide on its cell surface. In a

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preferred embodiment, the polynucleotides may be introduced using a viral expression system (e.g., vaccinia or other pox virus, retrovirus, or adenovirus), which may involve the use of a non-pathogenic (defective) virus. Techniques for incorporating polynucleotides into such expression systems are well known to those of ordinary skill in the art. The polynucleotides may also be administered as "naked" plasmid vectors as described, for example, in Ulmer et al., Science 259:1745-1749, 1993 and reviewed by Cohen, Science 259:1691-1692, 1993. Techniques for incorporating DNA into such vectors are well known to those of ordinary skill in the art. A retroviral vector may additionally transfer or incorporate a gene for a selectable marker (to aid in the identification or selection of transduced cells) and/or a targeting moiety, such as a gene that encodes a ligand for a receptor on a specific target cell, to render the vector target specific. Targeting may also be accomplished using an antibody, by methods known to those of ordinary skill in the art.

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Other formulations for therapeutic purposes include colloidal dispersion systems, such as macromolecule complexes, nanocapsules, microspheres, beads, and lipid-based systems including oil-in-water emulsions, micelles, mixed micelles, and liposomes. A preferred colloidal system for use as a delivery vehicle in vitro and in vivo is a liposome (i.e., an artificial membrane vesicle). The uptake of naked polynucleotides may be increased by incorporating the polynucleotides into and/or onto biodegradable beads, which are efficiently transported into the cells. The preparation and use of such systems is well known in the art.

In a related aspect, a polynucleotide vaccine as described above may be administered simultaneously with or sequentially to either a polypeptide of the present invention or a known *Chlamydia* antigen. For example, administration of polynucleotides encoding a polypeptide of the present invention, either "naked" or in a delivery system as described above, may be followed by administration of an antigen in order to enhance the protective immune effect of the vaccine.

Polypeptides and polynucleotides disclosed herein may also be employed in adoptive immunotherapy for the treatment of *Chlamydial* infection. Adoptive immunotherapy may be broadly classified into either active or passive immunotherapy. In active immunotherapy, treatment relies on the *in vivo* stimulation of the endogenous

host immune system with the administration of immune response-modifying agents (for example, vaccines, bacterial adjuvants, and/or cytokines).

In passive immunotherapy, treatment involves the delivery of biologic reagents with established immune reactivity (such as effector cells or antibodies) that 5 can directly or indirectly mediate anti-Chlamydia effects and does not necessarily depend on an intact host immune system. Examples of effector cells include T lymphocytes (for example, CD8+ cytotoxic T-lymphocyte, CD4+ T-helper), killer cells (such as Natural Killer cells, lymphokine-activated killer cells), B cells, or antigen presenting cells (such as dendritic cells and macrophages) expressing the disclosed antigens. The polypeptides disclosed herein may also be used to generate antibodies or anti-idiotypic antibodies (as in U.S. Patent No. 4,918,164), for passive immunotherapy.

The predominant method of procuring adequate numbers of T-cells for adoptive immunotherapy is to grow immune T-cells in vitro. Culture conditions for expanding single antigen-specific T-cells to several billion in number with retention of 15 antigen recognition in vivo are well known in the art. These in vitro culture conditions typically utilize intermittent stimulation with antigen, often in the presence of cytokines, such as IL-2, and non-dividing feeder cells. As noted above, the immunoreactive polypeptides described herein may be used to rapidly expand antigen-specific T cell cultures in order to generate sufficient number of cells for immunotherapy. In particular, antigen-presenting cells, such as dendritic, macrophage, monocyte, fibroblast, or B-cells, may be pulsed with immunoreactive polypeptides, or polynucleotide sequence(s) may be introduced into antigen presenting cells, using a variety of standard techniques well known in the art. For example, antigen presenting cells may be transfected or transduced with a polynucleotide sequence, wherein said 25 sequence contains a promoter region appropriate for increasing expression, and can be expressed as part of a recombinant virus or other expression system. Several viral vectors may be used to transduce an antigen presenting cell, including pox virus, vaccinia virus, and adenovirus; also, antigen presenting cells may be transfected with polynucleotide sequences disclosed herein by a variety of means, including gene-gun 30 technology, lipid-mediated delivery, electroporation, osmotic shock, and particlate delivery mechanisms, resulting in efficient and acceptable expression levels as determined by one of ordinary skill in the art. For cultured T-cells to be effective in

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therapy, the cultured T-cells must be able to grow and distribute widely and to survive long term in vivo. Studies have demonstrated that cultured T-cells can be induced to grow in vivo and to survive long term in substantial numbers by repeated stimulation with antigen supplemented with IL-2 (see, for example, Cheever, M., et al, "Therapy With Cultured T Cells: Principles Revisited," Immunological Reviews, 157:177, 1997).

The polypeptides disclosed herein may also be employed to generate and/or isolate chlamydial-reactive T-cells, which can then be administered to the patient. In one technique, antigen-specific T-cell lines may be generated by in vivo immunization with short peptides corresponding to immunogenic portions of the 10 disclosed polypeptides. The resulting antigen specific CD8+ or CD4+ T-cell clones may be isolated from the patient, expanded using standard tissue culture techniques, and returned to the patient.

Alternatively, peptides corresponding to immunogenic portions of the polypeptides may be employed to generate Chlamydia reactive T cell subsets by 15 selective in vitro stimulation and expansion of autologous T cells to provide antigenspecific T cells which may be subsequently transferred to the patient as described, for example, by Chang et al. (Crit. Rev. Oncol. Hematol., 22(3), 213, 1996). Cells of the immune system, such as T cells, may be isolated from the peripheral blood of a patient. using a commercially available cell separation system, such as IsolexTM System, available from Nexell Therapeutics, Inc. Irvine, CA. The separated cells are stimulated with one or more of the immunoreactive polypeptides contained within a delivery vehicle, such as a microsphere, to provide antigen-specific T cells. The population of antigen-specific T cells is then expanded using standard techniques and the cells are administered back to the patient.

In other embodiments, T-cell and/or antibody receptors specific for the polypeptides disclosed herein can be cloned, expanded, and transferred into other vectors or effector cells for use in adoptive immunotherapy. In particular, T cells may be transfected with the appropriate genes to express the variable domains from chlamydia specific monoclonal antibodies as the extracellular recognition elements and 30 joined to the T cell receptor signaling chains, resulting in T cell activation, specific lysis, and cytokine release. This enables the T cell to redirect its specificity in an MHCindependent manner. See for example, Eshhar, Z., Cancer Immunol Immunother, 45(3-

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4):131-6, 1997 and Hwu, P., et al, Cancer Res, 55(15):3369-73, 1995. Another embodiment may include the transfection of chlamydia antigen specific alpha and beta T cell receptor chains into alternate T cells, as in Cole, DJ, et al, Cancer Res, 55(4):748-52, 1995.

In a further embodiment, syngeneic or autologous dendritic cells may be pulsed with peptides corresponding to at least an immunogenic portion of a polypeptide disclosed herein. The resulting antigen-specific dendritic cells may either be transferred into a patient, or employed to stimulate T cells to provide antigen-specific T cells which may, in turn, be administered to a patient. The use of peptide-pulsed dendritic cells to generate antigen-specific T cells and the subsequent use of such antigen-specific T cells to eradicate disease in a murine model has been demonstrated by Cheever et al, Immunological Reviews, 157:177, 1997). Additionally, vectors expressing the disclosed polynucleotides may be introduced into stem cells taken from the patient and clonally propagated in vitro for autologous transplant back into the same patient.

15 Within certain aspects, polypeptides, polynucleotides, T cells and/or binding agents disclosed herein may be incorporated into pharmaceutical compositions or immunogenic compositions (i.e., vaccines). Alternatively, a pharmaceutical composition may comprise an antigen-presenting cell (e.g. a dendritic cell) transfected with a Chlamydial polynucleotide such that the antigen presenting cell expresses a Chlamydial polypeptide. Pharmaceutical compositions comprise one or more such compounds and a physiologically acceptable carrier. Vaccines may comprise one or more such compounds and an immunostimulant. An immunostimulant may be any substance that enhances or potentiates an immune response to an exogenous antigen. Examples of immunostimulants include adjuvants, biodegradable microspheres (e.g., 25 polylactic galactide) and liposomes (into which the compound is incorporated; see e.g., Fullerton, U.S. Patent No. 4,235,877). Vaccine preparation is generally described in. for example, M.F. Powell and M.J. Newman, eds., "Vaccine Design (the subunit and adjuvant approach)," Plenum Press (NY, 1995). Pharmaceutical compositions and vaccines within the scope of the present invention may also contain other compounds, which may be biologically active or inactive. For example, one or more immunogenic portions of other Chlamydial antigens may be present, either incorporated into a fusion polypeptide or as a separate compound, within the composition or vaccine.

A pharmaceutical composition or vaccine may contain DNA encoding one or more of the polypeptides as described above, such that the polypeptide is generated in situ. As noted above, the DNA may be present within any of a variety of delivery systems known to those of ordinary skill in the art, including nucleic acid expression systems, bacteria and viral expression systems. Numerous gene delivery techniques are well known in the art, such as those described by Rolland, Crit. Rev. Therap. Drug Carrier Systems 15:143-198, 1998, and references cited therein. Appropriate nucleic acid expression systems contain the necessary DNA sequences for expression in the patient (such as a suitable promoter and terminating signal). Bacterial delivery systems involve the administration of a bacterium (such as Bacillus-Calmette-Guerrin) that expresses an immunogenic portion of the polypeptide on its cell surface or secretes such an epitope.

In a preferred embodiment, the DNA may be introduced using a viral expression system (e.g., vaccinia or other pox virus, retrovirus, adenovirus, baculovirus, 15 togavirus, bacteriophage, and the like), which often involves the use of a non-pathogenic (defective), replication competent virus.

For example, many viral expression vectors are derived from viruses of the retroviridae family. This family includes the murine leukemia viruses, the mouse mammary tumor viruses, the human foamy viruses, Rous sarcoma virus, and the immunodeficiency viruses, including human, simian, and feline. Considerations when designing retroviral expression vectors are discussed in Comstock et al. (1997).

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Excellent murine leukemia virus (MLV)-based viral expression vectors have been developed by Kim et al. (1998). In creating the MLV vectors, Kim et al. found that the entire gag sequence, together with the immediate upstream region, could be deleted without significantly affecting viral packaging or gene expression. Further, it was found that nearly the entire U3 region could be replaced with the immediately-early promoter of human cytomegalovirus without deleterious effects. Additionally, MCR and internal ribosome entry sites (IRES) could be added without adverse effects. Based on their observations, Kim et al. have designed a series of MLV-based expression vectors comprising one or more of the features described above.

As more has been learned about human foamy virus (HFV), characteristics of HFV that are favorable for its use as an expression vector have been

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discovered. These characteristics include the expression of pol by splicing and start of translation at a defined initiation codon. Other aspects of HFV viral expression vectors are reviewed in Bodem et al. (1997).

Murakami et al. (1997) describe a Rous sarcoma virus (RSV)-based 5 replication-competent avian retrovirus vectors, IR1 and IR2 to express a heterologous gene at a high level. In these vectors, the IRES derived from encephalomyocarditis virus (EMCV) was inserted between the env gene and the heterologous gene. The IR1 vector retains the splice-acceptor site that is present downstream of the env gene while the IR2 vector lacks it. Murakami et al. have shown high level expression of several different heterologous genes by these vectors.

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Recently, a number of lentivirus-based retroviral expression vectors have been developed. Kafri et al. (1997) have shown sustained expression of genes delivered directly into liver and muscle by a human immunodeficiency virus (HIV)-based expression vector. One benefit of the system is the inherent ability of HIV to transduce non-dividing cells. Because the viruses of Kafri et al. are pseudotyped with vesicular stomatitis virus G glycoprotein (VSVG), they can transduce a broad range of tissues and cell types.

A large number of adenovirus-based expression vectors have been developed, primarily due to the advantages offered by these vectors in gene therapy applications. Adenovirus expression vectors and methods of using such vectors are the subject of a number of United States patents, including United States Patent No. 5,698,202, United States Patent No. 5,616,326, United States Patent No. 5,585,362, and United States Patent No. 5,518,913, all incorporated herein by reference.

Additional adenoviral constructs are described in Khatri et al. (1997) and 25 Tomanin et al. (1997). Khatri et al. describe novel ovine adenovirus expression vectors and their ability to infect bovine nasal turbinate and rabbit kidney cells as well as a range of human cell type, including lung and foreskin fibroblasts as well as liver, prostate, breast, colon and retinal lines. Tomanin et al. describe adenoviral expression vectors containing the T7 RNA polymerase gene. When introduced into cells 30 containing a heterologous gene operably linked to a T7 promoter, the vectors were able to drive gene expression from the T7 promoter. The authors suggest that this system may be useful for the cloning and expression of genes encoding cytotoxic proteins.

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Poxviruses are widely used for the expression of heterologous genes in mammalian cells. Over the years, the vectors have been improved to allow high expression of the heterologous gene and simplify the integration of multiple heterologous genes into a single molecule. In an effort to diminish cytopathic effects and to increase safety, vaccinia virus mutant and other poxviruses that undergo abortive infection in mammalian cells are receiving special attention (Oertli et al., 1997). The use of poxviruses as expression vectors is reviewed in Carroll and Moss (1997).

Togaviral expression vectors, which includes alphaviral expression vectors have been used to study the structure and function of proteins and for protein production purposes. Attractive features of togaviral expression vectors are rapid and efficient gene expression, wide host range, and RNA genomes (Huang, 1996). Also, recombinant vaccines based on alphaviral expression vectors have been shown to induce a strong humoral and cellular immune response with good immunological memory and protective effects (Tubulekas et al., 1997). Alphaviral expression vectors and their use are discussed, for example, in Lundstrom (1997).

In one study, Li and Garoff (1996) used Semliki Forest virus (SFV) expression vectors to express retroviral genes and to produce retroviral particles in BHK-21 cells. The particles produced by this method had protease and reverse transcriptase activity and were infectious. Furthermore, no helper virus could be detected in the virus stocks. Therefore, this system has features that are attractive for its use in gene therapy protocols.

Baculoviral expression vectors have traditionally been used to express heterologous proteins in insect cells. Examples of proteins include mammalian chemokine receptors (Wang et al., 1997), reporter proteins such as green fluorescent protein (Wu et al., 1997), and FLAG fusion proteins (Wu et al., 1997; Koh et al., 1997). Recent advances in baculoviral expression vector technology, including their use in virion display vectors and expression in mammalian cells is reviewed by Possee (1997). Other reviews on baculoviral expression vectors include Jones and Morikawa (1996) and O'Reilly (1997).

Other suitable viral expression systems are disclosed, for example, in Fisher-Hoch et al., *Proc. Natl. Acad. Sci. USA 86*:317-321, 1989; Flexner et al., *Ann. N.Y. Acad. Sci. 569*:86-103, 1989; Flexner et al., *Vaccine 8*:17-21, 1990; U.S. Patent

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Nos. 4,603,112, 4,769,330, and 5,017,487; WO 89/01973; U.S. Patent No. 4,777,127; GB 2,200,651; EP 0,345,242; WO 91/02805; Berkner, Biotechniques 6:616-627, 1988; Rosenfeld et al., Science 252:431-434, 1991; Kolls et al., Proc. Natl. Acad. Sci. USA 91:215-219, 1994; Kass-Eisler et al., Proc. Natl. Acad. Sci. USA 90:11498-11502, 1993; Guzman et al., Circulation 88:2838-2848, 1993; and Guzman et al., Circ. Res. 73:1202-1207, 1993. Techniques for incorporating DNA into such expression systems are well known to those of ordinary skill in the art. In other systems, the DNA may be introduced as "naked" DNA, as described, for example, in Ulmer et al., Science 259:1745-1749, 1993 and reviewed by Cohen, Science 259:1691-1692, 1993. The uptake of naked DNA may be increased by coating the DNA onto biodegradable beads, which are efficiently transported into the cells.

It will be apparent that a vaccine may comprise a polynucleotide and/or a polypeptide component, as desired. It will also be apparent that a vaccine may contain pharmaceutically acceptable salts of the polynucleotides and/or polypeptides provided herein. Such salts may be prepared from pharmaceutically acceptable non-toxic bases, including organic bases (e.g., salts of primary, secondary and tertiary amines and basic amino acids) and inorganic bases (e.g., sodium, potassium, lithium, ammonium, calcium and magnesium salts). While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, 20 the type of carrier will vary depending on the mode of administration. Compositions of the present invention may be formulated for any appropriate manner of administration, including for example, topical, oral, nasal, intravenous, intracranial, intraperitoneal, subcutaneous or intramuscular administration. For parenteral administration, such as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a fat, a 25 wax or a buffer. For oral administration, any of the above carriers or a solid carrier, such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactate polyglycolate) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4,897,268 and 5,075,109.

Such compositions may also comprise buffers (e.g., neutral buffered saline or phosphate buffered saline), carbohydrates (e.g., glucose, mannose, sucrose or dextrans), mannitol, proteins, polypeptides or amino acids such as glycine, antioxidants, bacteriostats, chelating agents such as EDTA or glutathione, adjuvants (e.g., aluminum hydroxide), solutes that render the formulation isotonic, hypotonic or weakly hypertonic with the blood of a recipient, suspending agents, thickening agents and/or preservatives.

5 Alternatively, compositions of the present invention may be formulated as a lyophilizate. Compounds may also be encapsulated within liposomes using well known technology.

Any of a variety of immunostimulants may be employed in the vaccines of this invention. For example, an adjuvant may be included. Most adjuvants contain a substance designed to protect the antigen from rapid catabolism, such as aluminum hydroxide or mineral oil, and a stimulator of immune responses, such as lipid A, Bortadella pertussis or Mycobacterium tuberculosis derived proteins. Suitable adjuvants are commercially available as, for example, Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, MI); Merck Adjuvant 65 (Merck and Company, Inc., Rahway, NJ); AS-2 (SmithKline Beecham, Philadelphia, PA); aluminum salts such as aluminum hydroxide gel (alum) or aluminum phosphate; salts of calcium, iron or zinc; an insoluble suspension of acylated tyrosine; acylated sugars; cationically or anionically derivatized polysaccharides; polyphosphazenes; biodegradable microspheres; monophosphoryl lipid A and quil A. Cytokines, such as GM-CSF or interleukin-2, -7, or -12, may also be used as adjuvants.

Within the vaccines provided herein, under select circumstances, the adjuvant composition may be designed to induce an immune response predominantly of the Th1 type or Th2 type. High levels of Th1-type cytokines (e.g., IFN-γ, TNFα, IL-2 and IL-12) tend to favor the induction of cell mediated immune responses to an administered antigen. In contrast, high levels of Th2-type cytokines (e.g., IL-4, IL-5, IL-6 and IL-10) tend to favor the induction of humoral immune responses. Following application of a vaccine as provided herein, a patient will support an immune response that includes Th1- and Th2-type responses. Within a preferred embodiment, in which a response is predominantly Th1-type, the level of Th1-type cytokines will increase to a greater extent than the level of Th2-type cytokines. The levels of these cytokines may be readily assessed using standard assays. For a review of the families of cytokines, see Mosmann and Coffman, Ann. Rev. Immunol. 7:145-173, 1989.

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Preferred adjuvants for use in eliciting a predominantly Th1-type response include, for example, a combination of monophosphoryl lipid A, preferably 3de-O-acylated monophosphoryl lipid A (3D-MPL), together with an aluminum salt. MPL adjuvants are available from Corixa Corporation (Seattle, WA; see US Patent Nos. 5 4,436,727; 4,877,611; 4,866,034 and 4,912,094). CpG-containing oligonucleotides (in which the CpG dinucleotide is unmethylated) also induce a predominantly Th1 response. Such oligonucleotides are well known and are described, for example, in WO 96/02555 and WO 99/33488. Immunostimulatory DNA sequences are also described, for example, by Sato et al., Science 273:352, 1996. Another preferred adjuvant is a saponin, preferably QS21 (Aquila Biopharmaceuticals Inc., Framingham, MA), which may be used alone or in combination with other adjuvants. For example, an enhanced system involves the combination of a monophosphoryl lipid A and saponin derivative, such as the combination of OS21 and 3D-MPL as described in WO 94/00153, or a less reactogenic composition where the OS21 is quenched with cholesterol, as described in WO 96/33739. Other preferred formulations comprise an oil-in-water emulsion and tocopherol. A particularly potent adjuvant formulation involving OS21, 3D-MPL and tocopherol in an oil-in-water emulsion is described in WO 95/17210.

Other preferred adjuvants include Montanide ISA 720 (Seppic, France), SAF (Chiron, California, United States), ISCOMS (CSL), MF-59 (Chiron), the SBAS 20 series of adjuvants (e.g., SBAS-2 or SBAS-4, available from SmithKline Beecham, Rixensart, Belgium), Detox (Corixa Corporation; Seattle, WA), RC-529 (Corixa Corporation; Seattle, WA) and other aminoalkyl glucosaminide 4-phosphates (AGPs), such as those described in pending U.S. Patent Application Serial Nos. 08/853,826 and 09/074.720, the disclosures of which are incorporated herein by reference in their entireties.

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Any vaccine provided herein may be prepared using well known methods that result in a combination of antigen, immunostimulant and a suitable carrier or excipient. The compositions described herein may be administered as part of a sustained release formulation (i.e., a formulation such as a capsule, sponge or gel (composed of polysaccharides, for example) that effects a slow release of compound following administration). Such formulations may generally be prepared using well known technology (see, e.g., Coombes et al., Vaccine 14:1429-1438, 1996) and

administered by, for example, oral, rectal or subcutaneous implantation, or by implantation at the desired target site. Sustained-release formulations may contain a polypeptide, polynucleotide or antibody dispersed in a carrier matrix and/or contained within a reservoir surrounded by a rate controlling membrane.

Carriers for use within such formulations are biocompatible, and may also be biodegradable; preferably the formulation provides a relatively constant level of active component release. Such carriers include microparticles of poly(lactide-coglycolide), as well as polyacrylate, latex, starch, cellulose and dextran. Other delayed-release carriers include supramolecular biovectors, which comprise a non-liquid hydrophilic core (e.g., a cross-linked polysaccharide or oligosaccharide) and, optionally, an external layer comprising an amphiphilic compound, such as a phospholipid (see e.g., U.S. Patent No. 5,151,254 and PCT applications WO 94/20078, WO/94/23701 and WO 96/06638). The amount of active compound contained within a sustained release formulation depends upon the site of implantation, the rate and expected duration of release and the nature of the condition to be treated or prevented.

Any of a variety of delivery vehicles may be employed within pharmaceutical compositions and vaccines to facilitate production of an antigen-specific immune response that targets *Chlamydia*-infected cells. Delivery vehicles include antigen presenting cells (APCs), such as dendritic cells, macrophages, B cells, monocytes and other cells that may be engineered to be efficient APCs. Such cells may, but need not, be genetically modified to increase the capacity for presenting the antigen, to improve activation and/or maintenance of the T cell response, to have anti-*Chlamydia* effects *per se* and/or to be immunologically compatible with the receiver (*i.e.*, matched HLA haplotype). APCs may generally be isolated from any of a variety of biological fluids and organs, and may be autologous, allogeneic, syngeneic or xenogeneic cells.

Certain preferred embodiments of the present invention use dendritic cells or progenitors thereof as antigen-presenting cells. Dendritic cells are highly potent APCs (Banchereau and Steinman, Nature 392:245-251, 1998) and have been shown to be effective as a physiological adjuvant for eliciting prophylactic or therapeutic immunity (see Timmerman and Levy, Ann. Rev. Med. 50:507-529, 1999). In general, dendritic cells may be identified based on their typical shape (stellate in situ, with

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marked cytoplasmic processes (dendrites) visible *in vitro*), their ability to take up, process and present antigens with high efficiency, and their ability to activate naïve T cell responses. Dendritic cells may, of course, be engineered to express specific cell-surface receptors or ligands that are not commonly found on dendritic cells *in vivo* or ex vivo, and such modified dendritic cells are contemplated by the present invention. As an alternative to dendritic cells, secreted vesicles antigen-loaded dendritic cells (called exosomes) may be used within a vaccine (see Zitvogel et al., Nature Med. 4:594-600, 1998).

Dendritic cells and progenitors may be obtained from peripheral blood, bone marrow, lymph nodes, spleen, skin, umbilical cord blood or any other suitable tissue or fluid. For example, dendritic cells may be differentiated ex vivo by adding a combination of cytokines such as GM-CSF, IL-4, IL-13 and/or TNFα to cultures of monocytes harvested from peripheral blood. Alternatively, CD34 positive cells harvested from peripheral blood, umbilical cord blood or bone marrow may be differentiated into dendritic cells by adding to the culture medium combinations of GM-CSF, IL-3, TNFα, CD40 ligand, LPS, flt3 ligand and/or other compound(s) that induce differentiation, maturation and proliferation of dendritic cells.

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Dendritic cells are conveniently categorized as "immature" and "mature" cells, which allows a simple way to discriminate between two well characterized phenotypes. However, this nomenclature should not be construed to exclude all possible intermediate stages of differentiation. Immature dendritic cells are characterized as APC with a high capacity for antigen uptake and processing, which correlates with the high expression of Fcy receptor and mannose receptor. The mature phenotype is typically characterized by a lower expression of these markers, but a high expression of cell surface molecules responsible for T cell activation such as class I and class II MHC, adhesion molecules (e.g., CD54 and CD11) and costimulatory molecules (e.g., CD40, CD80, CD86 and 4-1BB).

APCs may generally be transfected with a polynucleotide encoding a Chlamydial protein (or portion or other variant thereof) such that the Chlamydial polypeptide, or an immunogenic portion thereof, is expressed on the cell surface. Such transfection may take place ex vivo, and a composition or vaccine comprising such transfected cells may then be used for therapeutic purposes, as described herein. Alternatively, a gene delivery vehicle that targets a dendritic or other antigen presenting cell may be administered to a patient, resulting in transfection that occurs in vivo. In vivo and ex vivo transfection of dendritic cells, for example, may generally be performed using any methods known in the art, such as those described in WO 97/24447, or the gene gun approach described by Mahvi et al., Immunology and cell Biology 75:456-460, 1997. Antigen loading of dendritic cells may be achieved by incubating dendritic cells or progenitor cells with the Chlamydial polypeptide, DNA (naked or within a plasmid vector) or RNA; or with antigen-expressing recombinant bacterium or viruses (e.g., vaccinia, fowlpox, adenovirus or lentivirus vectors). Prior to loading, the polypeptide may be covalently conjugated to an immunological partner that provides T cell help (e.g., a carrier molecule). Alternatively, a dendritic cell may be pulsed with a non-conjugated immunological partner, separately or in the presence of the polypeptide.

Routes and frequency of administration of pharmaceutical compositions and vaccines, as well as dosage, will vary from individual to individual. In general, the pharmaceutical compositions and vaccines may be administered by injection (e.g., intracutaneous, intramuscular, intravenous or subcutaneous), intramasally (e.g., by aspiration) or orally. Between 1 and 3 doses may be administered for a 1-36 week period. Preferably, 3 doses are administered, at intervals of 3-4 months, and booster vaccinations may be given periodically thereafter. Alternate protocols may be appropriate for individual patients. A suitable dose is an amount of polypeptide or DNA that, when administered as described above, is capable of raising an immune response in an immunized patient sufficient to protect the patient from Chlamydial infection for at least 1-2 years. In general, the amount of polypeptide present in a dose (or produced in situ by the DNA in a dose) ranges from about 1 pg to about 10 pg to about 1 mg, and preferably from about 100 pg to about 1 mg, and preferably from about 100 pg to about 1 mg, suitable dose sizes will vary with the size of the patient, but will typically range from about 0.1 mL to about 5 mL.

While any suitable carrier known to those of ordinary skill in the art may
be employed in the pharmaceutical compositions of this invention, the type of carrier
will vary depending on the mode of administration. For parenteral administration, such
as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a fat, a
wax or a buffer. For oral administration, any of the above carriers or a solid carrier,

such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactic galactide) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable 5 microspheres are disclosed, for example, in U.S. Patent Nos, 4,897,268 and 5,075,109.

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In general, an appropriate dosage and treatment regimen provides the active compound(s) in an amount sufficient to provide therapeutic and/or prophylactic benefit. Such a response can be monitored by establishing an improved clinical outcome in treated patients as compared to non-treated patients. Increases in preexisting immune responses to a Chlamydial protein generally correlate with an improved clinical outcome. Such immune responses may generally be evaluated using standard proliferation, cytotoxicity or cytokine assays, which may be performed using samples obtained from a patient before and after treatment.

In another aspect, the present invention provides methods for using the polypeptides described above to diagnose Chlamydial infection. In this aspect, methods are provided for detecting Chlamydial infection in a biological sample, using one or more of the above polypeptides, either alone or in combination. For clarity, the term "polypeptide" will be used when describing specific embodiments of the inventive diagnostic methods. However, it will be clear to one of skill in the art that the fusion proteins of the present invention may also be employed in such methods.

As used herein, a "biological sample" is any antibody-containing sample obtained from a patient. Preferably, the sample is whole blood, sputum, serum, plasma, saliva, cerebrospinal fluid or urine. More preferably, the sample is a blood, serum or plasma sample obtained from a patient. The polypeptides are used in an assay, as described below, to determine the presence or absence of antibodies to the polypeptide(s) in the sample, relative to a predetermined cut-off value. The presence of such antibodies indicates previous sensitization to Chlamydia antigens which may be indicative of Chlamvdia-infection.

In embodiments in which more than one polypeptide is employed, the 30 polypeptides used are preferably complementary (i.e., one component polypeptide will tend to detect infection in samples where the infection would not be detected by another component polypeptide). Complementary polypeptides may generally be identified by

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using each polypeptide individually to evaluate serum samples obtained from a series of patients known to be infected with *Chlamydia*. After determining which samples test positive (as described below) with each polypeptide, combinations of two or more polypeptides may be formulated that are capable of detecting infection in most, or all, of the samples tested.

A variety of assay formats are known to those of ordinary skill in the art for using one or more polypeptides to detect antibodies in a sample. See, e.g., Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988, which is incorporated herein by reference. In a preferred embodiment, the assay involves the use of polypeptide immobilized on a solid support to bind to and remove the antibody from the sample. The bound antibody may then be detected using a detection reagent that contains a reporter group. Suitable detection reagents include antibodies that bind to the antibody/polypeptide complex and free polypeptide labeled with a reporter group (e.g., in a semi-competitive assay). Alternatively, a competitive assay may be utilized, in which an antibody that binds to the polypeptide is labeled with a reporter group and allowed to bind to the immobilized antigen after incubation of the antigen with the sample. The extent to which components of the sample inhibit the binding of the labeled antibody to the polypeptide is indicative of the reactivity of the sample with the immobilized polypeptide.

The solid support may be any solid material known to those of ordinary skill in the art to which the antigen may be attached. For example, the solid support may be a test well in a microtiter plate, or a nitrocellulose or other suitable membrane. Alternatively, the support may be a bead or disc, such as glass, fiberglass, latex or a plastic material such as polystyrene or polyvinylchloride. The support may also be a magnetic particle or a fiber optic sensor, such as those disclosed, for example, in U.S. Patent No. 5.359.681.

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The polypeptides may be bound to the solid support using a variety of techniques known to those of ordinary skill in the art. In the context of the present invention, the term "bound" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the antigen and functional groups on the support or may be a linkage by way of a cross-linking agent). Binding by adsorption to a well in a microtiter plate or to a membrane is preferred. In

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such cases, adsorption may be achieved by contacting the polypeptide, in a suitable buffer, with the solid support for a suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and 1 day. In general, contacting a well of a plastic microtiter plate (such as polystyrene or polyvinylchloride)

with an amount of polypeptide ranging from about 10 ng to about 1 µg, and preferably about 100 ng, is sufficient to bind an adequate amount of antigen.

Covalent attachment of polypeptide to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the polypeptide. For example, the polypeptide may be bound to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the polypeptide (see, e.g., Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

In certain embodiments, the assay is an enzyme linked immunosorbent
assay (ELISA). This assay may be performed by first contacting a polypeptide antigen
that has been immobilized on a solid support, commonly the well of a microtiter plate,
with the sample, such that antibodies to the polypeptide within the sample are allowed
to bind to the immobilized polypeptide. Unbound sample is then removed from the
immobilized polypeptide and a detection reagent capable of binding to the immobilized
antibody-polypeptide complex is added. The amount of detection reagent that remains
bound to the solid support is then determined using a method appropriate for the
specific detection reagent.

More specifically, once the polypeptide is immobilized on the support as described above, the remaining protein binding sites on the support are typically blocked. Any suitable blocking agent known to those of ordinary skill in the art, such as bovine serum albumin (BSA) or Tween 20™ (Sigma Chemical Co., St. Louis, MO) may be employed. The immobilized polypeptide is then incubated with the sample, and antibody is allowed to bind to the antigen. The sample may be diluted with a suitable dilutent, such as phosphate-buffered saline (PBS) prior to incubation. In general, an appropriate contact time (i.e., incubation time) is that period of time that is sufficient to detect the presence of antibody within an HGE-infected sample. Preferably, the contact time is sufficient to achieve a level of binding that is at least 95% of that achieved at

equilibrium between bound and unbound antibody. Those of ordinary skill in the art will recognize that the time necessary to achieve equilibrium may be readily determined by assaying the level of binding that occurs over a period of time. At room temperature, an incubation time of about 30 minutes is generally sufficient.

Unbound sample may then be removed by washing the solid support with an appropriate buffer, such as PBS containing 0.1% Tween 20™. Detection reagent may then be added to the solid support. An appropriate detection reagent is any compound that binds to the immobilized antibody-polypeptide complex and that can be detected by any of a variety of means known to those in the art. Preferably, the detection reagent contains a binding agent (such as, for example, Protein A, Protein G, immunoglobulin, lectin or free antigen) conjugated to a reporter group. Preferred reporter groups include enzymes (such as horseradish peroxidase), substrates, cofactors, inhibitors, dyes, radionuclides, luminescent groups, fluorescent groups and biotin. The conjugation of binding agent to reporter group may be achieved using standard methods 15 known to those of ordinary skill in the art. Common binding agents may also be purchased conjugated to a variety of reporter groups from many commercial sources (e.g., Zymed Laboratories, San Francisco, CA, and Pierce, Rockford, IL).

The detection reagent is then incubated with the immobilized antibodypolypeptide complex for an amount of time sufficient to detect the bound antibody. An appropriate amount of time may generally be determined from the manufacturer's instructions or by assaying the level of binding that occurs over a period of time. Unbound detection reagent is then removed and bound detection reagent is detected using the reporter group. The method employed for detecting the reporter group depends upon the nature of the reporter group. For radioactive groups, scintillation 25 counting or autoradiographic methods are generally appropriate. Spectroscopic methods may be used to detect dyes, luminescent groups and fluorescent groups. Biotin may be detected using avidin, coupled to a different reporter group (commonly a radioactive or fluorescent group or an enzyme). Enzyme reporter groups may generally be detected by the addition of substrate (generally for a specific period of time), followed by spectroscopic or other analysis of the reaction products.

To determine the presence or absence of anti-Chlamydia antibodies in the sample, the signal detected from the reporter group that remains bound to the solid

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support is generally compared to a signal that corresponds to a predetermined cut-off value. In one preferred embodiment, the cut-off value is the average mean signal obtained when the immobilized antigen is incubated with samples from an uninfected patient. In general, a sample generating a signal that is three standard deviations above the predetermined cut-off value is considered positive for Chlamydia-infection. In an alternate preferred embodiment, the cut-off value is determined using a Receiver Operator Curve, according to the method of Sackett et al., Clinical Epidemiology: A Basic Science for Clinical Medicine, Little Brown and Co., 1985, pp. 106-107. Briefly, in this embodiment, the cut-off value may be determined from a plot of pairs of true positive rates (i.e., sensitivity) and false positive rates (100%-specificity) that correspond to each possible cut-off value for the diagnostic test result. The cut-off value on the plot that is the closest to the upper left-hand corner (i.e., the value that encloses the largest area) is the most accurate cut-off value, and a sample generating a signal that is higher than the cut-off value determined by this method may be considered positive. Alternatively, the cut-off value may be shifted to the left along the plot, to minimize the false positive rate, or to the right, to minimize the false negative rate. In general, a sample generating a signal that is higher than the cut-off value determined by this method is considered positive for Chlamydial infection.

In a related embodiment, the assay is performed in a rapid flow-through or strip test format, wherein the antigen is immobilized on a membrane, such as 20 nitrocellulose. In the flow-through test, antibodies within the sample bind to the immobilized polypeptide as the sample passes through the membrane. A detection reagent (e.g., protein A-colloidal gold) then binds to the antibody-polypeptide complex as the solution containing the detection reagent flows through the membrane. The detection of bound detection reagent may then be performed as described above. In the 25 strip test format, one end of the membrane to which polypeptide is bound is immersed in a solution containing the sample. The sample migrates along the membrane through a region containing detection reagent and to the area of immobilized polypeptide. Concentration of detection reagent at the polypeptide indicates the presence of anti-Chlamydia antibodies in the sample. Typically, the concentration of detection reagent at that site generates a pattern, such as a line, that can be read visually. The absence of such a pattern indicates a negative result. In general, the amount of polypeptide

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immobilized on the membrane is selected to generate a visually discernible pattern when the biological sample contains a level of antibodies that would be sufficient to generate a positive signal in an ELISA, as discussed above. Preferably, the amount of polypeptide immobilized on the membrane ranges from about 25 ng to about 1 µg, and more preferably from about 50 ng to about 500 ng. Such tests can typically be performed with a very small amount (e.g., one drop) of patient serum or blood.

Of course, numerous other assay protocols exist that are suitable for use with the polypeptides of the present invention. The above descriptions are intended to be exemplary only. One example of an alternative assay protocol which may be usefully employed in such methods is a Western blot, wherein the proteins present in a biological sample are separated on a gel, prior to exposure to a binding agent. Such techniques are well known to those of skill in the art.

The present invention further provides agents, such as antibodies and antigen-binding fragments thereof, that specifically bind to a Chlamydial protein. As used herein, an antibody, or antigen-binding fragment thereof, is said to "specifically bind" to a Chlamydial protein if it reacts at a detectable level (within, for example, an ELISA) with a Chlamydial protein, and does not react detectably with unrelated proteins under similar conditions. As used herein, "binding" refers to a noncovalent association between two separate molecules such that a complex is formed. The ability to bind may be evaluated by, for example, determining a binding constant for the formation of the complex. The binding constant is the value obtained when the concentration of the complex is divided by the product of the component concentrations. In general, two compounds are said to "bind," in the context of the present invention, when the binding constant for complex formation exceeds about 10³ L/mol. The binding constant may be determined using methods well known in the art.

Binding agents may be further capable of differentiating between patients with and without a *Chlamydial* infection using the representative assays provided herein. In other words, antibodies or other binding agents that bind to a *Chlamydial* protein will generate a signal indicating the presence of a *Chlamydial* infection in at least about 20% of patients with the disease, and will generate a negative signal indicating the absence of the disease in at least about 90% of individuals without infection. To determine whether a binding agent satisfies this requirement, biological

samples (e.g., blood, sera, sputum urine and/or tissue biopsies) from patients with and without Chlamydial infection (as determined using standard clinical tests) may be assayed as described herein for the presence of polypeptides that bind to the binding agent. It will be apparent that a statistically significant number of samples with and 5 without the disease should be assayed. Each binding agent should satisfy the above criteria; however, those of ordinary skill in the art will recognize that binding agents may be used in combination to improve sensitivity.

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Any agent that satisfies the above requirements may be a binding agent. For example, a binding agent may be a ribosome, with or without a pertide component, an RNA molecule or a polypeptide. In a preferred embodiment, a binding agent is an antibody or an antigen-binding fragment thereof. Antibodies may be prepared by any of a variety of techniques known to those of ordinary skill in the art. See, e.g., Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988. In general, antibodies can be produced by cell culture techniques, including the generation of monoclonal antibodies as described herein, or via transfection of antibody genes into suitable bacterial or mammalian cell hosts, in order to allow for the production of recombinant antibodies. In one technique, an immunogen comprising the polypeptide is initially injected into any of a wide variety of mammals (e.g., mice, rats, rabbits, sheep or goats). In this step, the polypeptides of this invention may serve as the immunogen without modification. Alternatively, particularly for relatively short polypeptides, a superior immune response may be elicited if the polypeptide is joined to a carrier protein, such as bovine serum albumin or keyhole limpet hemocyanin. The immunogen is injected into the animal host, preferably according to a predetermined schedule incorporating one or more booster immunizations, and the animals are bled periodically. 25 Polyclonal antibodies specific for the polypeptide may then be purified from such antisera by, for example, affinity chromatography using the polypeptide coupled to a suitable solid support.

Monoclonal antibodies specific for an antigenic polypeptide of interest may be prepared, for example, using the technique of Kohler and Milstein, Eur. J. Immunol. 6:511-519, 1976, and improvements thereto. Briefly, these methods involve the preparation of immortal cell lines capable of producing antibodies having the desired specificity (i.e., reactivity with the polypeptide of interest). Such cell lines may WO 02/08267

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be produced, for example, from spleen cells obtained from an animal immunized as described above. The spleen cells are then immortalized by, for example, fusion with a myeloma cell fusion partner, preferably one that is syngeneic with the immunized animal. A variety of fusion techniques may be employed. For example, the spleen cells and myeloma cells may be combined with a nonionic detergent for a few minutes and then plated at low density on a selective medium that supports the growth of hybrid cells, but not myeloma cells. A preferred selection technique uses HAT (hypoxanthine, aminopterin, thymidine) selection. After a sufficient time, usually about 1 to 2 weeks, colonies of hybrids are observed. Single colonies are selected and their culture supernatants tested for binding activity against the polypeptide. Hybridomas having high reactivity and specificity are preferred.

Monoclonal antibodies may be isolated from the supernatants of growing hybridoma colonies. In addition, various techniques may be employed to enhance the yield, such as injection of the hybridoma cell line into the peritoneal cavity of a suitable vertebrate host, such as a mouse. Monoclonal antibodies may then be harvested from the ascites fluid or the blood. Contaminants may be removed from the antibodies by conventional techniques, such as chromatography, gel filtration, precipitation, and extraction. The polypeptides of this invention may be used in the purification process in, for example, an affinity chromatography step.

Within certain embodiments, the use of antigen-binding fragments of antibodies may be preferred. Such fragments include Fab fragments, which may be prepared using standard techniques. Briefly, immunoglobulins may be purified from rabbit serum by affinity chromatography on Protein A bead columns (Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, 1988) and digested by papain to yield Fab and Fc fragments. The Fab and Fc fragments may be separated by affinity chromatography on protein A bead columns.

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Monoclonal antibodies of the present invention may be coupled to one or more therapeutic agents. Suitable agents in this regard include radionuclides, differentiation inducers, drugs, toxins, and derivatives thereof. Preferred radionuclides include ⁹⁰Y, ¹²³I, ¹²⁵I, ¹³¹I, ¹⁸⁶Re, ¹⁸⁸Re, ²¹¹At, and ²¹²Bi. Preferred drugs include methotrexate, and pyrimidine and purine analogs. Preferred differentiation inducers include phorbol esters and butyric acid. Preferred toxins include ricin, abrin, diptheria

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toxin, cholera toxin, gelonin, Pseudomonas exotoxin, Shigella toxin, and pokeweed antiviral protein.

A therapeutic agent may be coupled (e.g., covalently bonded) to a suitable monoclonal antibody either directly or indirectly (e.g., via a linker group). A 5 direct reaction between an agent and an antibody is possible when each possesses a substituent capable of reacting with the other. For example, a nucleophilic group, such as an amino or sulfhydryl group, on one may be capable of reacting with a carbonylcontaining group, such as an anhydride or an acid halide, or with an alkyl group containing a good leaving group (e.g., a halide) on the other.

Alternatively, it may be desirable to couple a therapeutic agent and an antibody via a linker group. A linker group can function as a spacer to distance an antibody from an agent in order to avoid interference with binding capabilities. A linker group can also serve to increase the chemical reactivity of a substituent on an agent or an antibody, and thus increase the coupling efficiency. An increase in chemical reactivity may also facilitate the use of agents, or functional groups on agents, which otherwise would not be possible.

It will be evident to those skilled in the art that a variety of bifunctional or polyfunctional reagents, both homo- and hetero-functional (such as those described in the catalog of the Pierce Chemical Co., Rockford, IL), may be employed as the linker group. Coupling may be effected, for example, through amino groups, carboxyl groups, sulfhydryl groups or oxidized carbohydrate residues. There are numerous references describing such methodology, e.g., U.S. Patent No. 4,671,958, to Rodwell et al.

Where a therapeutic agent is more potent when free from the antibody portion of the immunoconjugates of the present invention, it may be desirable to use a linker group which is cleavable during or upon internalization into a cell. A number of different cleavable linker groups have been described. The mechanisms for the intracellular release of an agent from these linker groups include cleavage by reduction of a disulfide bond (e.g., U.S. Patent No. 4,489,710, to Spitler), by irradiation of a photolabile bond (e.g., U.S. Patent No. 4.625,014, to Senter et al.), by hydrolysis of derivatized amino acid side chains (e.g., U.S. Patent No. 4,638,045, to Kohn et al.), by serum complement-mediated hydrolysis (e.g., U.S. Patent No. 4,671,958, to Rodwell et al.), and acid-catalyzed hydrolysis (e.g., U.S. Patent No. 4,569,789, to Blattler et al.).

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It may be desirable to couple more than one agent to an antibody. In one embodiment, multiple molecules of an agent are coupled to one antibody molecule. In another embodiment, more than one type of agent may be coupled to one antibody. Regardless of the particular embodiment, immunoconjugates with more than one agent 5 may be prepared in a variety of ways. For example, more than one agent may be coupled directly to an antibody molecule, or linkers which provide multiple sites for attachment can be used. Alternatively, a carrier can be used.

A carrier may bear the agents in a variety of ways, including covalent bonding either directly or via a linker group. Suitable carriers include proteins such as albumins (e.g., U.S. Patent No. 4,507,234, to Kato et al.), peptides and polysaccharides such as aminodextran (e.g., U.S. Patent No. 4,699,784, to Shih et al.). A carrier may also bear an agent by noncovalent bonding or by encapsulation, such as within a liposome vesicle (e.g., U.S. Patent Nos. 4,429,008 and 4,873,088). Carriers specific for radionuclide agents include radiohalogenated small molecules and chelating 15 compounds. For example, U.S. Patent No. 4,735,792 discloses representative radiohalogenated small molecules and their synthesis. A radionuclide chelate may be formed from chelating compounds that include those containing nitrogen and sulfur atoms as the donor atoms for binding the metal, or metal oxide, radionuclide. For example, U.S. Patent No. 4,673,562, to Davison et al. discloses representative chelating compounds and their synthesis.

A variety of routes of administration for the antibodies and immunoconjugates may be used. Typically, administration will be intravenous, intramuscular, subcutaneous or in site-specific regions by appropriate methods. It will be evident that the precise dose of the antibody/immunoconjugate will vary depending upon the antibody used, the antigen density, and the rate of clearance of the antibody.

Antibodies may be used in diagnostic tests to detect the presence of Chlamydia antigens using assays similar to those detailed above and other techniques well known to those of skill in the art, thereby providing a method for detecting Chlamydial infection in a patient.

Diagnostic reagents of the present invention may also comprise DNA sequences encoding one or more of the above polypeptides, or one or more portions thereof. For example, at least two oligonucleotide primers may be employed in a

polymerase chain reaction (PCR) based assay to amplify Chlamydia-specific cDNA derived from a biological sample, wherein at least one of the oligonucleotide primers is specific for a DNA molecule encoding a polypeptide of the present invention. The presence of the amplified cDNA is then detected using techniques well known in the art, such as gel electrophoresis. Similarly, oligonucleotide probes specific for a DNA molecule encoding a polypeptide of the present invention may be used in a hybridization assay to detect the presence of an inventive polypeptide in a biological sample.

As used herein, the term "oligonucleotide primer/probe specific for a

DNA molecule" means an oligonucleotide sequence that has at least about 80%,
preferably at least about 90% and more preferably at least about 95%, identity to the

DNA molecule in question. Oligonucleotide primers and/or probes which may be
usefully employed in the inventive diagnostic methods preferably have at least about
10-40 nucleotides. In a preferred embodiment, the oligonucleotide primers comprise at
1s least about 10 contiguous nucleotides of a DNA molecule encoding one of the
polypeptides disclosed herein. Preferably, oligonucleotide probes for use in the
inventive diagnostic methods comprise at least about 15 contiguous oligonucleotides of
a DNA molecule encoding one of the polypeptides disclosed herein. Techniques for
both PCR based assays and hybridization assays are well known in the art (see, for
example, Mullis et al. Ibid; Ehrlich, Ibid). Primers or probes may thus be used to detect
Chlamydia-specific sequences in biological samples. DNA probes or primers
comprising oligonucleotide sequences described above may be used alone or in
combination with each other.

The following Examples are offered by way of illustration and not by 25 way of limitation.

EXAMPLE 1

ISOLATION OF DNA SEQUENCES ENCODING CHLAMYDIA ANTIGENS

Chlamydia antigens of the present invention were isolated by expression cloning of a genomic DNA library of Chlamydia trachomatis LGV II essentially as described by Sanderson et al. (J. Exp. Med., 1995, 182:1751-1757) and were shown to induce PBMC proliferation and IFN-y in an immunoreactive T cell line.

A Chlamydia-specific T cell line was generated by stimulating PBMCs from a normal donor with no history of chlamydial genital tract infection with elementary bodies of Chlamydia trachomatis LGV II. This T cell line, referred to as TCL-8, was found to recognize both Chlamydia trachomatis and Chlamydia pneumonia 5 infected monocyte-derived dendritic cells.

A randomly sheared genomic library of Chlamydia trachomatis LGV II was constructed in Lambda ZAP (Stratagene, La Jolla, CA) and the amplified library plated out in 96 well microtiter plates at a density of 30 clones/well. Bacteria were induced to express recombinant protein in the presence of 2 mM IPTG for 3 h, then 10 pelleted and resuspended in 200 µl of RPMI 10% FBS. 10 µl of the induced bacterial suspension was transferred to 96 well plates containing autologous monocyte-derived dendritic cells. After a 2 h incubation, dendritic cells were washed to remove free E. coli and Chlamydia-specific T cells were added. Positive E. coli pools were identified by determining IFN-y production and proliferation of the T cells in response to the 15 pools.

Four positive pools were identified, which were broken down to yield four pure clones (referred to as 1-B1-66, 4-D7-28, 3-G3-10 and 10-C10-31), with insert sizes of 481 bp, 183 bp, 110 bp and 1400 bp, respectively. The determined DNA sequences for 1-B1-66, 4-D7-28, 3-G3-10 and 10-C10-31 are provided in SEO ID NO: 1-4, respectively. Clone 1-B1-66 is approximately in region 536690 of the C. trachomatis genome (NCBI C. trachomatis database). Within clone 1-B1-66, an open reading frame (ORF) has been identified (nucleotides 115 - 375) that encodes a previously identified 9 kDa protein (Stephens, et al. Genbank Accession No. AE001320), the sequence of which is provided in SEQ ID NO: 5). Clone 4-D7-28 is a smaller region of the same ORF (amino acids 22-82 of 1-B1-66). Clone 3-G3-10 is 25 approximately in region 74559 of the C. trachomatis genome. The insert is cloned in the antisense orientation with respect to its orientation in the genome. The clone 10-C10-31 contains an open reading frame that corresponds to a previously published sequence for S13 ribosomal protein from Chlamydia trachomatis (Gu, L. et al. J. 30 Bacteriology, 177:2594-2601, 1995). The predicted protein sequences for 4-D7-28 and 10-C10-31 are provided in SEQ ID NO: 6 and 12, respectively. Predicted protein sequences for 3-G3-10 are provided in SEQ ID NO: 7-11.

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In a related series of screening studies, an additional T cell line was used to screen the genomic DNA library of Chlamydia trachomatis LGV II described above. A Chlamydia-specific T cell line (TCT-1) was derived from a patient with a chlamydial genital tract infection by stimulating patient PBMC with autologous monocyte-derived dendritic cells infected with elementary bodies of Chlamydia trachomatis LGV II. One clone, 4C9-18 (SEQ ID NO: 21), containing a 1256 bp insert, elicited a specific immune response, as measured by standard proliferation assays, from the Chlamydia-specific T cell line TCT-1. Subsequent analysis revealed this clone to contain three known sequences: lipoamide dehydrogenase (Genbank Accession No. AE001326), disclosed in SEQ ID NO: 22; a hypothetical protein CT429 (Genbank Accession No. AE001316), disclosed in SEQ ID NO: 23; and part of an open reading frame of ubiquinone methyltransferase CT428 (Genbank Accession No. AE001316), disclosed in SEQ ID NO: 24.

In further studies involving clone 4C9-18 (SEQ ID NO: 21), the fulllength amino acid sequence for lipoamide dehydrognase (SEQ ID NO: 22) from C. trachomatis (LGV II) was expressed in clone CtL2-LPDA-FL, as disclosed in SEQ ID NO: 90.

To further characterize the open reading frame containing the T cell stimulating epitope(s), a cDNA fragment containing nucleotides 1-695 of clone 4C9-18 with a cDNA sequence encoding a 6X-Histidine tag on the amino terminus was subcloned into the Ndel/EcoRl site of the pET17b vector (Novagen, Madison, WI), referred to as clone 4C9-18#2 BL21 pLysS (SEQ ID NO: 25, with the corresponding amino acid sequence provided in SEQ ID NO: 26) and transformed into E. coli. Selective induction of the transformed E. coli with 2 mM IPTG for three hours resulted in the expression of a 26 kDa protein from clone 4C9-18#2 BL21 pLysS, as evidenced by standard Coomassie-stained SDS-PAGE. To determine the immunogenicity of the protein encoded by clone 4C9-18#2 BL21 pLysS, E. coli expressing the 26 kDa protein were titered onto 1 x 10⁴ monocyte-derived dendritic cells and incubated for two hours. The dendritic cell cultures were washed and 2.5 x 10⁴ T cells (TCT-1) added and allowed to incubate for an additional 72 hours, at which time the level of IFN-7 in the culture supernatant was determined by ELISA. As shown in Fig. 1, the T-cell line TCT-1 was found to respond to induced cultures as measured by IFN-e, indicating a

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**Chlamydia-specific T-cell response against the lipoamide dehydrogenase sequence.

Similarly, the protein encoded by clone 4C9-18#2 BL21 pLysS was shown to stimulate.

the TCT-1 T-cell line by standard proliferation assays.

Subsequent studies to identify additional Chlamydia trachomatis

5 antigens using the above-described CD4+ T-cell expression cloning technique yielded
additional clones. The TCT-1 and TCL-8 Chlamydia-specific T-cell lines, as well as
the TCP-21 T-cell line were utilized to screen the Chlamydia trachomatis LGVII
genomic library. The TCP-21 T-cell line was derived from a patient having a humoral
immune response to Chlamydia pruemoniae. The TCT-1 cell line identified 37 positive

10 pools, the TCT-3 cell line identified 41 positive pools and the TCP-21 cell line
identified 2 positive pools. The following clones were derived from 10 of these positive
pools. Clone 11-A3-93 (SEQ ID NO: 64), identified by the TCP-21 cell line, is a 1339
bp genomic fragment sharing homology to the HAD superfamily (CT103). The second
insert in the same clone shares homology with the fab I gene (CT104) present on the
15 complementary strand. Clone 11-C12-91 (SEQ ID NO: 63), identified using the TCP21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares

homology with the 60 kDa cysteine rich outer membrane protein of C. pnuemoniae.

Clone 11-G10-46, (SEQ ID NO: 62), identified using the TCT-3 cell line, contains a 688 bp insert that shares homology to the hypothetical protein CT610.

20 Clone 11-G1-34, (SEQ ID NO: 61), identified using the TCT-3 cell line, has two partial open reading frames (ORF) with an insert size of 1215 bp. One ORF shares homology to the malate dehydrogenase gene (CT376), and the other ORF shares homology to the glycogen hydrolase gene (CT042). Clone 11-H3-68, (SEQ ID NO: 60), identified using the TCT-3 cell line, has two ORFs with a total insert size of 1180 bp. One partial ORF encodes the plasmid-encoded PGP6-D virulence protein while the second ORF is a complete ORF for the L1 ribosomal gene (CT318). Clone 11-H4-28, (SEQ ID NO: 59), identified using the TCT-3 cell line, has an insert size of 552 bp and is part of the ORF for the dnaK gene (CT396). Clone 12-B3-95, (SEQ ID NO: 58), identified using the TCT-1 cell line, has an insert size of 463 bp and is a part of the ORF for for the lipoamide dehydrogenase gene (CT557). Clones 15-G1-89 and 12-B3-95 are identical, (SEQ ID NO: 55 and 58, respectively), identified using the TCT-1 cell line, has an insert size of 463 bp and is part of the incompile dehydrogenase gene

(CT557). Clone 12-G3-83, (SEQ ID NO: 57), identified using the TCT-1 cell line, has an insert size of 1537 bp and has part of the ORF for the hypothetical protein CT622.

Clone 23-G7-68, (SEQ ID NO: 79), identified using the TCT-3 cell line, contains a 950 bp insert and contains a small part of the L11 ribosomal ORF, the entire 5 ORF for L1 ribosomal protein and a part of the ORF for L10 ribosomal protein. In addition, this clone also identified the patient lines CT4, CT5, CT11, CT12, and CHH037. Clone 22-F8-91. (SEO ID NO: 80), identified using the TCT-1 cell line. contains a 395 bp insert that contains a part of the pmpC ORF on the complementary strand of the clone. Clone 21-E8-95, (SEQ ID NO: 81), identified using the TCT-3 cell 10 line, contains a 2,085 bp insert which contains part of CT613 ORF, the complete ORF for CT612, the complete ORF for CT611 and part of the ORF for CT610. Clone 19-F12-57, (SEQ ID NO: 82), identified using the TCT-3 cell line, contains a 405 bp insert which contains part of the CT 858 ORF and a small part of the recA ORF. Clone 19-F12-53, (SEO ID NO; 83), identified using the TCT-3 cell line, contains a 379 bp insert 15 that is part of the ORF for CT455 encoding glutamyl tRNA synthetase. Clone 19-A5-54, (SEO ID NO: 84), identified using the TCT-3 cell line, contains a 715 bp insert that is part of the ORF3 (complementary strand of the clone) of the cryptic plasmid. Clone 17-E11-72, (SEQ ID NO: 85), identified using the TCT-1 cell line, contains a 476 bp insert that is part of the ORF for Opp 2 and pmpD. The pmpD region of this clone is 20 covered by the pmpD region of clone 15-H2-76. Clone 17-C1-77, (SEO ID NO: 86), identified using the the patient cell lines CT3, CT1, CT4, and CT12, contains a 1551 bp insert that is part of the CT857 ORF, as well as part of the CT858 ORF. Clone 15-H2-76, (SEQ ID NO: 87), identified using the TCT-1 cell line, contains a 3,031 bp insert that contains a large part of the pmpD ORF, part of the CT089 ORF, as well as part of 25 the ORF for SvcE. Clone 15-A3-26, (SEO ID NO: 88), contains a 976 bp insert that contains part of the ORF for CT858. Clone 17-G4-36, (SEQ ID NO: 267), identified using the patient lines CL8, TCT-10, CT1, CT5, CT13, and CHH037, contains a 680 bp insert that is in frame with beta-gal in the plasmid and shares homology to part of the ORF for DNA-directed RNA polymerase beta subunit (CT315 in SerD).

Several of the clones described above share homology to various polymorphic membrane proteins. The genomic sequence of *Chlamydia trachomatis* contains a family of nine polymorphic membrane protein genes, referred to as pmp. These genes are designated pmpA, pmpB, pmpC, pmpD, pmpE, pmpF, pmpG, pmpH and pmpI. Proteins expressed from these genes are believed to be of biological relevance in generating a protective immune response to a *Chlamydial* infection. In particular, pmpC, pmpD, pmpE and pmpI contain predictable signal peptides, suggesting they are outer membrane proteins, and therefore, potential immunological targets.

Based on the Chlamydia trachomatis LGVII serovar sequence, primer pairs were designed to PCR amplify the full-length fragments of pmpC, pmpD, pmpE, pmpG, pmpH and pmpI. The resulting fragments were subcloned into the DNA vaccine 10 vector JA4304 or JAL, which is JA4304 with a modified linker (SmithKline Beecham, London, England). Specifically, PmpC was subcloned into the JAL vector using the 5' oligo GAT AGG CGC GCC GCA ATC ATG AAA TTT ATG TCA GCT ACT GCT G and the 3' oligo CAG AAC GCG TTT AGA ATG TCA TAC GAG CAC CGC A, as provided in SEQ ID NO: 197 and 198, respectively. PCR amplification of the gene 15 under conditions well known in the art and ligation into the 5' ASCI/3' MluI sites of the JAL vector was completed after inserting the short nucleotide sequence GCAATC (SEO ID NO: 199) upstream of the ATG to create a Kozak-like sequence. The resulting expression vector contained the full-length pmpC gene comprising 5325 nucleotides (SEO ID NO: 173) containing the hypothetical signal sequence, which encodes a 187 20 kD protein (SEO ID NO: 179). The pmpD gene was subcloned into the JA4304 vaccine vector following PCR amplification of the gene using the following oligos: 5' oligo-TGC AAT CAT GAG TTC GCA GAA AGA TAT AAA AAG C (SEQ ID NO: 200) and 3' oligo- CAG AGC TAG CTT AAA AGA TCA ATC GCA ATC CAG TAT TC (SEQ ID NO: 201). The gene was ligated into the a 5' blunted HIII/3' MluI site of the 25 JA4304 vaccine vector using standard techniques well known in the art. The CAATC (SEO ID NO: 202) was inserted upstream of the ATG to create a Kozak-like sequence. This clone is unique in that the last threonine of the HindIII site is missing due to the blunting procedure, as is the last glycine of the Kozak-like sequence. The insert, a 4593 nucleotide fragment (SEQ ID NO: 172) is the full-length gene for pmpD containing the hypothetical signal sequence, which encodes a 161 kD protein (SEO ID NO: 178). PmpE was subcloned into the JA4304 vector using the 5' oligo- TGC AAT CAT GAA AAA AGC GTT TTT CTT TTT C (SEQ ID NO: 203), and the 3' oligo- CAG AAC

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GCG TCT AGA ATC GCA GAG CAA TTT C (SEQ ID NO: 204). Following PCR amplification, the gene was ligated into the 5' blunted HIII/3' MluI site of JA4304. To facilitate this, a short nucleotide sequence, TGCAATC (SEQ ID NO; 293), was added upstream of the initiation codon for creating a Kozak-like sequence and reconstituting 5 the HindIII site. The insert is the full-length pmpE gene (SEQ ID NO: 171) containing the hypothetical signal sequence. The pmpE gene encodes a 105 kD protein (SEO ID NO: 177). The pmpG gene was PCR amplified using the 5' oligo- GTG CAA TCA TGA TTC CTC AAG GAA TTT ACG (SEQ ID NO: 205), and the 3' oligo- CAG AAC GCG TTT AGA ACC GGA CTT TAC TTC C (SEQ ID NO: 206) and subcloned 10 into the JA4304 vector. Similar cloning strategies were followed for the pmpI and pmpK genes. In addition, primer pairs were designed to PCR amplify the full-length or overlapping fragments of the pmp genes, which were then subcloned for protein expression in the pET17b vector (Novagen, Madison, WI) and transfected into E. coli BL21 pLysS for expression and subsequent purification utilizing the histidine-nickel chromatographic methodology provided by Novagen. Several of the genes encoding the recombinant proteins, as described below, lack the native signal sequence to facilitate expression of the protein. Full-length protein expression of pmpC was accomplished through expression of two overlapping fragments, representing the amino and carboxy termini. Subcloning of the pmpC-amino terminal portion, which lacks the signal sequence, (SEQ ID NO: 187, with the corresponding amino acid sequence provided in SEO ID NO: 195) used the 5' oligo- CAG ACA TAT GCA TCA CCA TCA CCA TCA CGA GGC GAG CTC GAT CCA AGA TC (SEQ ID NO: 207), and the 3' oligo- CAG AGG TAC CTC AGA TAG CAC TCT CTC CTA TTA AAG TAG G (SEQ ID NO: 208) into the 5' NdeI/3' KPN cloning site of the vector. The carboxy terminus portion 25 of the gene, pmpC-carboxy terminal fragment (SEQ ID NO: 186, with the corresponding amino acid sequence provided in SEQ ID NO: 194), was subcloned into the 5' Nhel/3' KPN cloning site of the expression vector using the following primers: 5' oligo- CAG AGC TAG CAT GCA TCA CCA TCA CCA TCA CGT TAA GAT TGA GAA CTT CTC TGG C (SEO ID NO: 209), and 3' oligo- CAG AGG TAC CTT AGA ATG TCA TAC GAG CAC CGC AG (SEQ ID NO: 210). PmpD was also expressed as two overlapping proteins. The pmpD-amino terminal portion, which lacks the signal sequence, (SEO ID NO: 185, with the corresponding amino acid sequence provided in

SEQ ID NO: 193) contains the initiating codon of the pET17b and is expressed as a 80 kD protein. For protein expression and purification purposes, a six-histidine tag follows the initiation codon and is fused at the 28th amino acid (nucleotide 84) of the gene. The following primers were used, 5' oligo, CAG ACA TAT GCA TCA CCA TCA CCA 5 TCA CGG GTT AGC (SEO ID NO: 211), and the 3' oligo- CAG AGG TAC CTC AGC TCC TCC AGC ACA CTC TCT TC (SEQ ID NO: 212), to splice into the 5' NdeI/3' KPN cloning site of the vector. The pmpD-carboxy terminus portion (SEO ID NO: 184) was expressed as a 92 kD protein (SEO ID NO: 192). For expression and subsequent purification, an additional methionine, alanine and serine was included, which represent the initiation codon and the first two amino acids from the pET17b vector. A six-histidine tag downstream of the methionine, alanine and serine is fused at the 691st amino acid (nucleotide 2073) of the gene. The 5' oligo- CAG AGC TAG CCA TCA CCA TCA CCA TCA CGG TGC TAT TTC TTG CTT ACG TGG (SEO ID. NO: 213) and the 3' oligo- CAG AGG TAC TTn AAA AGA TCA ATC GCA ATC 15 CAG TAT TCG (SEO ID NO: 214) were used to subclone the insert into the 5' NheI/3' KPN cloning site of the expression vector. PmpE was expressed as a 106kD protein (SEQ ID NO: 183 with the corresponding amino acid sequence provided in SEO ID NO: 191). The pmpE insert also lacks the native signal sequence. PCR amplification of the gene under conditions well known in the art was performed using the following 20 oligo primers: 5' oligo- CAG AGG ATC CAC ATC ACC ATC ACC ATC ACG GAC TAG CTA GAG AGG TTC (SEQ ID NO: 215), and the 3' oligo- CAG AGA ATT CCT AGA ATC GCA GAG CAA TTT C (SEO ID NO: 216), and the amplified insert was ligated into a 5' BamHI/3' EcoRI site of JA4304. The short nucleotide sequence, as provided in SEQ ID NO: 217, was inserted upstream of the initiation codon for creating 25 the Kozak-like sequence and reconstituting the HindIII site. The expressed protein contains the initiation codon and the downstream 21 amino acids from the pET17b expression vector, i.e., MASMTGGQQMGRDSSLVPSSDP (SEQ ID NO: 218). In addition, a six-histidine tag is included upstream of the sequence described above and is fused at the 28th amino acid (nucleotide 84) of the gene, which eliminates the hypothetical signal peptide. The sequences provided in SEO ID NO: 183 with the 30 corresponding amino acid sequence provided in SEQ ID NO: 191 do not include these additional sequences. The pmpG gene (SEQ ID NO: 182, with the corresponding

amino acid sequence provided in SEQ ID No; 190) was PCR amplified under conditions well known in the art using the following oligo primers: 5' oligo- CAG AGG TAC CGC ATC ACC ATC ACC ATC ACA TGA TTC CTC AAG GAA TTT ACG (SEQ ID NO: 219), and the 3' oligo- CAG AGC GGC CGC TTA GAA CCG GAC 5 TTT ACT TCC (SEO ID NO: 220), and ligated into the 5' KPN/3' NotI cloning site of the expression vector. The expressed protein contains an additional amino acid sequence at the amino end, namely, MASMTGGQQNGRDSSLVPHHHHHHH (SEQ ID NO: 221), which comprises the initiation codon and additional sequence from the pET17b expression vector. The pmpI gene (SEO ID NO: 181, with the corresponding 10 amino acid sequence provided in SEQ ID No; 189) was PCR amplified under conditions well known in the art using the following oligo primers: 5' oligo- CAG AGC TAG CCA TCA CCA TCA CCA TCA CCT CTT TGG CCA GGA TCC C (SEO ID NO: 222), and the 3' oligo- CAG AAC TAG TCT AGA ACC TGT AAG TGG TCC (SEQ ID NO: 223), and ligted into the expression vector at the 5' NheI/3' SpeI cloning site. The 95 kD expressed protein contains the initiation codon plus an additional alanine and serine from the pET17b vector at the amino end of the protein. In addition, a six-histidine tag is fused at the 21st amino acid of the gene, which eliminates the hypothetical signal peptide.

Clone 14H1-4, (SEQ ID NO: 56), identified using the TCT-3 cell line,
contains a complete ORF for the TSA gene, thiol specific antioxidant – CT603 (the
CT603 ORF is a homolog of CPn0778 from *C. pnuemoniae*). The TSA open reading
frame in clone 14-H1-4 was amplified such that the expressed protein possess an
additional methionine and a 6x histidine tag (amino terminal end). This amplified insert
was sub-cloned into the Nde/EcoRI sites of the pET17b vector. Upon induction of this
clone with IPTG, a 22.6 kDa protein was purified by Ni-NTA agarose affinity
chromatography. The determined amino acid sequence for the 195 amino acid ORF of
clone 14-H1-4 encoding the TSA gene is provided in SEQ ID NO: 65. Further analysis
yielded a full-length clone for the TSA gene, referred to as CTL2-TSA-FL, with the
full-length amino acid sequence provided in SEQ ID NO: 92.

Further studies yielded 10 additional clones identified by the TCT-1 and TCT-3 T-cell lines, as described above. The clones identified by the TCT-1 line are: 16-D4-22, 17-C5-19, 18-C5-2, 20-G3-45 and 21-C7-66; clones identified by the TCT-3

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cell line are: 17-C10-31, 17-E2-9, 22-A1-49 and 22-B3-53. Clone 21-G12-60 was recognized by both the TCT-1 and TCT-3 T cell lines. In addition, clone 20-G3-45, which contained sequence specific for pmpB, was identified against the patient lines CT1 and CT4. Clone 16-D4-22 (SEO ID NO: 119), identified using the TCT-1 cell line 5 contains a 953 bp insert that contains two genes, parts of open reading frame 3 (ORF3) and ORF4 of the C. trachomatis plasmid for growth within mammalian cells. Clone 17-C5-19 (SEO ID NO: 118), contains a 951 bp insert that contains part of the ORF for DT431, encoding for clpP 1 protease and part of the ORF for CT430 (diaminopimelate epimerase). Clone 18-C5-2 (SEQ ID NO: 117) is part of the ORF for S1 ribosomal 10 protein with a 446 bp insert that was identified using the TCT-1 cell line. Clone 20-G3-45 (SEQ ID NO: 116), identified by the TCT-1 cell line, contains a 437 bp insert that is part of the pmpB gene (CT413). Clone 21-C7-8 (SEQ ID NO: 115), identified by the TCT-1 line, contains a 995bp insert that encodes part of the dnaK like protein. The insert of this clone does not overlap with the insert of the TCT-3 clone 11-H4-28 (SEO 15 ID NO: 59), which was shown to be part of the dnaK gene CT396. Clone 17-C10-31 (SEQ ID NO: 114), identified by the TCT-3 cell line, contains a 976 bp insert. This clone contains part of the ORF for CT858, a protease containing IRBP and DHR domains. Clone 17-E2-9 (SEQ ID NO: 113) contains part of ORFs for two genes, CT611 and CT610, that span a 1142 bp insert. Clone 22-A1-49 (SEQ ID NO: 112), identified using the TCT-3 line, also contains two genes in a 698 bp insert. Part of the ORF for CT660 (DNA gyrase{gyrA 2}) is present on the top strand where as the complete ORF for a hypothetical protein CT659 is present on the complementary strand. Clone 22-B3-53 (SEQ ID NO: 111), identified by the TCT-1 line, has a 267 bp insert that encodes part of the ORF for GroEL (CT110). Clone 21-G12-60 (SEO ID 25 NO: 110), identified by both the TCT-1 and TCT-3 cell lines contains a 1461 bp insert that contains partial ORFs for hypothetical proteins CT875, CT229 and CT228.

Additional Chlamydia antigens were obtained by screening a genomic expression library of Chlamydia trachomatis (LGV II scrovar) in Lambda Screen-1 vector (Novagen, Madison, WI) with sera pooled from several Chlamydia-infected individuals using techniques well known in the art. The following immuno-reactive clones were identified and the inserts containing Chlamydia genes sequenced: CTL2#1 (SEQ ID NO: 71); CTL2#2 (SEQ ID NO: 70); CTL2#3-5' (SEQ ID NO: 72, a first

determined genomic sequence representing the 5' end); CTL2#3-3' (SEO ID NO: 73, a second determined genomic sequence representing the 3' end); CTL2#4 (SEQ ID NO: 53); CTL2#5 (SEQ ID NO: 69); CTL2#6 (SEQ ID NO: 68); CTL2#7 (SEQ ID NO: 67); CTL2#8b (SEQ ID NO: 54); CTL2#9 (SEQ ID NO: 66); CTL2#10-5' (SEQ ID NO: 74. 5 a first determined genomic sequence representing the 5' end); CTL2#10-3' (SEO ID NO: 75, a second determined genomic sequence representing the 3' end); CTL2#11-5' (SEO ID NO: 45, a first determined genomic sequence representing the 5' end); CTL2#11-3' (SEO ID NO: 44, a second determined genomic sequence representing the 3' end); CTL2#12 (SEQ ID NO: 46); CTL2#16-5' (SEQ ID NO: 47); CTL2#18-5' 10 (SEQ ID NO: 49, a first determined genomic sequence representing the 5' end); CTL2#18-3' (SEO ID NO: 48, a second determined genomic sequence representing the 3' end); CTL2#19-5' (SEO ID NO: 76, the determined genomic sequence representing the 5' end); CTL2#21 (SEQ ID NO: 50); CTL2#23 (SEQ ID NO: 51; and CTL2#24 (SEO ID NO: 52).

Additional Chlamydia trachomatis antigens were identified by 15 serological expression cloning. These studies used sera pooled from several Chlamydia-infected individuals, as described above, but, IgA, and IgM antibodies were used in addition to IgG as a secondary antibody. Clones screened by this method enhance detection of antigens recognized by an early immune response to a Chlamvdial 20 infection, that is a mucosal humoral immune response. The following immunoreactive clones were characterized and the inserts containing Chlamydia genes sequenced: CTL2gam-1 (SEO ID NO: 290), CTL2gam-2 (SEO ID NO: 289), CTL2gam-5 (SEO ID NO: 288), CTL2gam-6-3' (SEQ ID NO: 287, a second determined genomic sequence representing the 3' end), CTL2gam-6-5' (SEQ ID NO: 286, a first determined genomic 25 sequence representing the 5' end), CTL2gam-8 (SEQ ID NO: 285), CTL2gam-10 (SEQ ID NO: 284), CTL2gam-13 (SEQ ID NO: 283), CTL2gam-15-3' (SEQ ID NO: 282, a second determined genomic sequence representing the 3' end), CTL2gam-15-5' (SEQ ID NO: 281, a first determined genomic sequence representing the 5' end), CTL2gam-17 (SEO ID NO: 280), CTL2gam-18 (SEO ID NO: 279), CTL2gam-21 (SEQ ID NO: 30 278), CTL2gam-23 (SEO ID NO: 277), CTL2gam-24 (SEO ID NO: 276), CTL2gam-26 (SEQ ID NO: 275), CTL2gam-27 (SEQ ID NO: 274), CTL2gam-28 (SEQ ID NO: 273), CTL2gam-30-3' (SEQ ID NO: 272, a second determined genomic sequence

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representing the 3' end) and CTL2gam-30-5' (SEO ID NO: 271, a first determined genomic sequence representing the 5' end).

EXAMPLE 2

INDUCTION OF T CELL PROLIFERATION AND INTERFERON-Y PRODUCTION BY CHLAMYDIA TRACHOMATIS ANTIGENS

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The ability of recombinant Chlamydia trachomatis antigens to induce T cell proliferation and interferon-v production is determined as follows.

Proteins are induced by IPTG and purified by Ni-NTA agarose affinity chromatograph (Webb et al., J. Immunology 157:5034-5041, 1996). The purified polypeptides are then screened for the ability to induce T-cell proliferation in PBMC preparations. PBMCs from C. trachomatis patients as well as from normal donors whose T-cells are known to proliferate in response to Chlamvdia antigens, are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 µg/ml gentamicin. Purified polypeptides are added in duplicate at concentrations of 0.5 to 10 µg/mL. After six days of culture in 96-well round-bottom plates in a volume of 200 µl, 50 µl of medium is removed from each well for determination of IFN-y levels, as described below. The plates are then pulsed with 1 uCi/well of tritiated 20 thymidine for a further 18 hours, harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

IFN-y is measured using an enzyme-linked immunosorbent assay (ELISA). ELISA plates are coated with a mouse monoclonal antibody directed to human IFN-y (PharMingen, San Diego, CA) in PBS for four hours at room temperature. Wells are then blocked with PBS containing 5% (W/V) non-fat dried milk for 1 hour at room temperature. The plates are washed six times in PBS/0.2% TWEEN-20 and samples diluted 1:2 in culture medium in the ELISA plates are incubated overnight at 30 room temperature. The plates are again washed and a polyclonal rabbit anti-human IFN-y serum diluted 1:3000 in PBS/10% normal goat serum is added to each well. The plates are then incubated for two hours at room temperature, washed and horseradish peroxidase-coupled anti-rabbit IgG (Sigma Chemical So., St. Louis, MO) is added at a 1:2000 dilution in PBS/5% non-fat dried milk. After a further two hour incubation at room temperature, the plates are washed and TMB substrate added. The reaction is stopped after 20 min with 1 N sulfuric acid. Optical density is determined at 450 nm using 570 nm as a reference wavelength. Fractions that result in both replicates giving an OD two fold greater than the mean OD from cells cultured in medium alone, plus 3 standard deviations, are considered positive.

Using the above methodology, recombinant 1B1-66 protein (SEQ ID NO: 5) as well as two synthetic peptides corresponding to amino acid residues 48-67 (SEQ ID NO: 13; referred to as 1-B1-66/48-67) and 58-77 (SEQ ID NO: 14, referred to as 1B1-66/58-77), respectively, of SEQ ID NO: 5, were found to induce a proliferative response and IFN-γ production in a Chlamydia-specific T cell line used to screen a genomic library of C. trachomatis LGV II.

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Further studies have identified a *C. trachomatis*-specific T-cell epitope in the ribosomal S13 protein. Employing standard epitope mapping techniques well known in the art, two T-cell epitopes in the ribosomal S13 protein (rS13) were identified with a *Chlamydla*-specific T-cell line from donor CL-8 (T-cell line TCL-8 EB/DC). Fig. 8 illustrates that the first peptide, rS13 1-20 (SEQ ID NO: 106), is 100% identical with the corresponding *C. pneumoniae* sequence, explaining the cross-reactivity of the T-cell line to recombinant *C. trachomatis*- and *C. pneumoniae*-rS13. The response to the second peptide rS13 56-75 (SEQ ID NO: 108) is *C. trachomatis*-specific, indicating that the rS13 response in this healthy asymptomatic donor was elicited by exposure to *C. trachomatis* and not to *C. pneumoniae*, or any other microbial infection.

As described in Example 1, Clone 11-C12-91 (SEQ ID NO: 63), identified using the TCP-21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares homology with the 60 kDa cysteine rich outer membrane protein of *C. pneumoniae*, referred to as OMCB. To further define the reactive epitope(s), epitope mapping was performed using a series of overlapping peptides and the immunoassay previously described. Briefly, proliferative responses were determined by stimulating 2.5 x 10⁴ TCP-21 T-cells in the presence of 1 x 10⁴ monocyte-derived dendritic cells

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with either non-infectious elementary bodies derived from C. trachomatis and C. pneumoniae, or peptides derived from the protein sequence of C. 'trachomatis or C. pneumoniae OMCB protein (0.1 µg/ml). The TCP-21 T-cells responded to epitopes CT-OMCB #167-186, CT-OMCB #171-190, CT-OMCB #171-186, and to a lesser extent, CT-OMCB #175-186 (SEQ ID NO: 249-252, respectively). Notably, the TCP-21 T-cell line also gave a proliferative response to the homologous C. pneumoniae peptide CP-OMCB #171-186 (SEQ ID NO: 253), which was equal to or greater than the response to the C. trachomatis peptides. The amino acid substitutions in position two (i.e., Asp for Glu) and position four (i.e., Cys for Ser) did not alter the proliferative response of the T-cells and therefore demonstrating this epitope to be a cross-reactive epitope between C. trachomatis and C. pneumoniae.

To further define the epitope described above, an additional T-cell line, TCT-3, was used in epitope mapping experiments. The immunoassays were performed as described above, except that only peptides from *C. trachomatis* were tested. The T-cells gave a proliferative response to two peptides, CT-OMCB #152-171 and CT-OMCB #157-176 (SEQ ID NO: 246 and 247, respectively), thereby defining an additional immunogenic epitope in the cysteine rich outer membrane protein of *C. trachomatis*.

Clone 14H1-4, (SEQ ID NO: 56, with the corresponding full-length amino acid sequence provided in SEQ ID NO: 92), was identified using the TCT-3 cell line in the CD4 T-cell expression cloning system previously described, and was shown to contain a complete ORF for the, thiol specific antioxidant gene (CT603), referred to as TSA. Epitope mapping immunoassays were performed, as described above, to further define the epitope. The TCT-3 T-cells line exhibited a strong proliferative response to the overlapping peptides CT-TSA #96-115, CT-TSA #101-120 and CT-TSA #106-125 (SEQ ID NO: 254-256, respectively) demonstrating an immunoreactive epitope in the thiol specific antioxidant gene of *C. trachomatis* serovar LGVII.

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EXAMPLE 3 PREPARATION OF SYNTHETIC POLYPEPTIDES

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Polypeptides may be synthesized on a Millipore 9050 peptide synthesizer using FMOC chemistry with HPTU (O-Benzotriazole-N.N.N',N'-tetramethyluronium hexafluorophosphate) activation. A Gly-Cys-Gly sequence may be attached to the amino terminus of the peptide to provide a method of conjugating or labeling of the peptide. Cleavage of the peptides from the solid support may be carried out using the following cleavage mixture: trifluoroacetic acid:ethanedithiol:thioanisole:water:phenol (40:1:2:2:3). After cleaving for 2 hours, the peptides may be precipitated in cold methyl-t-butyl-ether. The peptide pellets may then be dissolved in water containing 0.1% trifluoroacetic acid (TFA) and Ivophilized prior to purification by C18 reverse phase HPLC. A gradient of 0-60% acetonitrile (containing 0.1% TFA) in water 15 (containing 0.1% TFA) may be used to elute the peptides. Following lyophilization of the pure fractions, the peptides may be characterized using electrospray mass spectrometry and by amino acid analysis.

EXAMPLE 4

ISOLATION AND CHARACTERIZATION OF DNA SEOUENCES ENCODING CHLAMYDIA ANTIGENS USING RETROVIRAL EXPRESSION VECTOR SYSTEMS AND SUBSEQUENT IMMUNOLOGICAL ANALYSIS

A genomic library of Chlamydia trachomatis LGV II was constructed by limited digests using BamHI, BglII, BstYi and MboI restriction enzymes. The restriction digest fragments were subsequently ligated into the BamHI site of the retroviral vectors pBIB-KS1,2,3. This vector set was modified to contain a Kosak translation initiation site and stop codons in order to allow expression of proteins from short DNA genomic fragments, as shown in Fig. 2. DNA pools of 80 clones were 30 prepared and transfected into the retroviral packaging line Phoenix-Ampho, as described in Pear, W.S., Scott, M.L. and Nolan, G.P., Generation of High Titre, Helperfree Retroviruses by Transient Transfection. Methods in Molecular Medicine: Gene

Therapy Protocols, Humana Press, Totowa, NJ, pp. 41-57. The Chlamydia library in retroviral form was then transduced into H2-Ld expressing P815 cells, which were then used as target cells to stimulate an antigen specific T-cell line.

A Chlamydia-specific, murine H2d restricted CD8+ T-cell line was 5 expanded in culture by repeated rounds of stimulation with irradiated C. trachomatisinfected J774 cells and irradiated syngeneic spleen cells, as described by Starnbach, M., in J. Immunol., 153:5183, 1994. This Chlamvdia-specific T-cell line was used to screen the above Chlamydia genomic library expressed by the retrovirally-transduced P815 cells. Positive DNA pools were identified by detection of IFN-y production using Elispot analysis (SEE Lalvani et al., J. Experimental Medicine 186:859-865, 1997).

Two positive pools, referred to as 2C7 and 2E10, were identified by IFNy Elispot assays. Stable transductants of P815 cells from pool 2C7 were cloned by limiting dilution and individual clones were selected based upon their capacity to elicit IFN-y production from the Chlamydia-specific CTL line. From this screening process, 15 four positive clones were selected, referred to as 2C7-8, 2C7-9, 2C7-19 and 2C7-21. Similarly, the positive pool 2E10 was further screened, resulting in an additional positive clone, which contains three inserts. The three inserts are fragments of the CT016, tRNA syntase and clpX genes (SEO ID NO: 268-270, respectively).

Transgenic DNA from these four positive 2C7 clones were PCR amplified using pBIB-KS specific primers to selectively amplify the Chlamydia DNA insert. Amplified inserts were gel purified and sequenced. One immunoreactive clone, 2C7-8 (SEO ID NO: 15, with the predicted amino acid sequence provided in SEO ID NO: 32), is a 160 bp fragment with homology to nucleotides 597304-597145 of Chlamydia trachomatis, serovar D (NCBI, BLASTN search; SEQ ID NO: 33, with the 25 predicted amino acid sequence provided in SEQ ID NO: 34). The sequence of clone 2C7-8 maps within two putative open reading frames from the region of high homology described immediately above, and in particular, one of these putative open reading frames, consisting of a 298 amino acid fragment (SEQ ID NO: 16, with the predicted amino acid sequence provided in SEO ID NO: 17), was demonstrated to exhibit immunological activity.

Full-length cloning of the 298 amino acid fragment (referred to as CT529 and/or the Cap1 gene) from serovar L2 was obtained by PCR amplification using 5'-

ttttgaagcaggtaggtgatattg (forward) (SEQ ID NO: 159) and 5'-ttaagaaatttaaaaaatccctta (reverse) (SEQ ID NO: 160) primers, using purified *C. trachomatis* L2 genomic DNA as template. This PCR product was gel-purified, cloned into pCRBlunt (Invitrogen, Carlsbad, CA) for sequencing, and then subcloned into the *EcoRI* site of pBIB-KMS, a derivative of pBIB-KS for expression. The *Chlamydia pnuemoniae* homlogue of CT529 is provided in SEQ ID NO: 291, with the corresponding amino acid sequence provided in SEQ ID NO: 292.

Full-length DNA encoding various CT529 serovars were amplified by PCR from bacterial lysates containing 105 IFU, essentially as described (Denamur, E., C. Sayada, A. Souriau, J. Orfila, A. Rodolakis and J. Elion. 1991. J. Gen. Microbiol. 10 137: 2525). The following serovars were amplified as described: Ba (SEO ID NO: 134, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 135); E (BOUR) and E (MTW447) (SEO ID NO: 122, with the corresponding predicted amino acid sequence provided in SEO ID NO: 123); F (NI1) (SEO ID NO: 128, with the corresponding predicted amino acid sequence provided in SEO ID NO: 129); G: (SEO ID NO: 126, with the corresponding predicted amino acid sequence provided in SEO ID NO: 127); Ia (SEQ ID NO: 124, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 125); L1 (SEQ ID NO: 130, with the corresponding predicted amino acid sequence provided in SEO ID NO: 131); L3 (SEO ID NO: 132, with the 20 corresponding predicted amino acid sequence provided in SEQ ID NO: 133); I (SEQ ID NO: 263, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 264); K (SEQ ID NO: 265, with the corresponding predicted amino acid sequence provided in SEO ID NO: 266); and MoPn (SEO ID NO: 136, with the corresponding predicted amino acid sequence provided in SEQ ID NO: 137). PCR reactions were performed with Advantage Genomic PCR Kit (Clontech, Palo Alto, CA) using primers 25 specific for serovar L2 DNA (external to the ORF). Primers sequences were 5'ggtataatatctctctaaattttg (forward-SEQ ID NO: 161) and 5'-agataaaaaaggctgtttc' (reverse-SEQ ID NO: 162) except for MoPn which required 5'-ttttgaagcaggtaggtgaatatg (forward-SEQ ID NO: 163) and 5'-tttacaataagaaaagctaagcactttgt (reverse-SEQ ID NO: 164), PCR amplified DNA was purified with OIAquick PCR purification kit (Oiagen, Valencia, CA) and cloned in pCR2.1 (Invitrogen, Carlsbad, CA) for sequencing.

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Sequencing of DNA derived from PCR amplified inserts of immunoreactive clones was done on an automated sequencer (ABI 377) using both a pBIB-KS specific forward primer 5'-cettacacagtectgetgac (SEQ ID NO: 165) and a reverse primer 3'-gtttccgggccctcacattg (SEQ ID NO: 166). PCRBlunt cloned DNA coding for CT529 serovar L2 and pCR2.1 cloned DNA coding for CT529 serovar Ba, E (BOUR), E (MTW447), F (NI1), G, Ia, K, L1, L3 and MoPn were sequenced using T7 promoter primer and universal M13 forward and M13 reverse primers.

To determine if these two putative open reading frames (SEO ID NO: 16 and 20) encoded a protein with an associated immunological function, overlapping 10 peptides (17-20 amino acid lengths) spanning the lengths of the two open reading frames were synthesized, as described in Example 3. A standard chromium release assay was utilized to determine the percent specific lysis of peptide-pulsed H2d restricted target cells. In this assay, aliquots of P815 cells (H2d) were labeled at 37° C for one hour with 100 μCi of 51Cr in the presence or absence of 1 μg/ml of the indicated peptides. Following this incubation, labeled P815 cells were washed to remove excess 51Cr and peptide, and subsequently plated in duplicate in microculture plates at a concentration of 1,000 cells/well. Effector CTL (Chlamydia-specific CD8 T cells) were added at the indicated effector:target ratios. Following a 4 hour incubation, supernatants were harvested and measured by gamma-counter for release of 51Cr into 20 the supernatant. Two overlapping peptides from the 298 amino acid open reading frame did specifically stimulate the CTL line. The peptides represented in SEQ ID NO: 138-156 were synthesized, representing the translation of the L2 homologue of the serovar D open reading frame for CT529 (Cap1 gene) and 216 amino acid open reading frame. As shown in Fig. 3, peptides CtC7.8-12 (SEO ID NO: 18, also referred to as Cap1#132-147, SEQ ID NO: 139) and CtC7.8-13 (SEQ ID NO: 19, also referred to as Cap1#138-155, SEO ID NO: 140) were able to elicit 38 to 52% specific lysis, respectively, at an effector to target ratio of 10:1. Notably, the overlap between these two peptides contained a predicted H2^d (K^d and L^d) binding peptide. A 10 amino acid peptide was synthesized to correspond to this overlapping sequence (SEQ ID NO: 31) and was 30 found to generate a strong immune response from the anti-Chlamvdia CTL line by elispot assay. Significantly, a search of the most recent Genbank database revealed no proteins have previously been described for this gene. Therefore, the putative open

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reading frame encoding clone 2C7-8 (SEO ID NO: 15) defines a gene which encompasses an antigen from Chlamvdia capable of stimulating antigen-specific CD8+ T-cells in a MHC-I restricted manner, demonstrating this antigen could be used to develop a vaccine against Chlamydia.

To confirm these results and to further map the epitope, truncated peptides (SEO ID NO: 138-156) were made and tested for recognition by the T-cells in an IFN-g ELISPOT assay. Truncations of either Ser139 (Cap1#140-147, SEQ ID NO: 146) or Leu147 (Cap1#138-146, SEO ID NO: 147) abrogate T-cell recognition. These results indicate that the 9-mer peptide Cap1#139-147 (SFIGGITYL, SEQ ID NO: 145) is the minimal epitope recognized by the Chlamydia-specific T-cells.

Sequence alignments of Cap1 (CT529) from selected serovars of C. trachomatis (SEO ID NO: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139) shows one of the amino acid differences is found in position 2 of the proposed epitope. The homologous serovar D peptide is SIIGGITYL (SEQ ID NO: 168). The ability of 15 SFIGGITYL and SIIGGITYL to target cells for recognition by the Chlamydia specific T-cells was compared. Serial dilutions of each peptide were incubated with P815 cells and tested for recognition by the T-cells in a 51 Cr release assay, as described above. The Chlamydia-specific T-cells recognize the serovar L2 peptide at a minimum concentration of 1 nM and the serovar D peptide at a minimum concentration of 10 nM.

Further studies have shown that a Cap1#139-147-specific T-cell clone recognizes C. trachomatis infected cells. To confirm that Caplus is presented on the surface of Chlamydia infected cells, Balb-3T3 (H-2d) cells were infected with C. trachomatis serovar L2 and tested to determine whether these cells are recognized by a CD8+ T-cell clone specific for Cap1#139-147 epitope (SEQ ID NO: 145). The T-cell 25 clone specific for Cap1#139-147 epitope was obtained by limiting dilution of the line 69 T-cells. The T-cell clone specifically recognized the Chlamydia infected cells. In these experiments, target cells were C. trachomatis infected (positive control) or uninfected Balb/3T3 cells, showing 45%, 36% and 30% specific lysis at 30:1, 10:1 and 3:1 effector to target ratios, respectively; or Cap1#139-147 epitope (SEQ ID NO: 145) 30 coated, or untreated P815 cells, showing 83%, 75% and 58% specific lysis at 30:1, 10:1 and 3:1 effector to target ratios, respectively (negative controls having less than 5% lysis in all cases). This data suggests that the epitope is presented during infection.

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In vivo studies show Cap1#139-147 epitope-specific T-cells are primed during murine infection with C. trachomatis. To determine if infection with C. trachomatis primes a Cap1#139-147 epitope-specific T-cell response, mice were infected i.p. with 108 IFU of C. trachomatis serovar L2. Two weeks after infection, the 5 mice were sacrificed and spleen cells were stimulated on irradiated syngeneic spleen cells pulsed with Cap1#139-147 epitope peptide. After 5 days of stimulation, the cultures were used in a standard 51Cr release assay to determine if there were Cap1#139-147 epitope-specific T-cells present in the culture. Specifically, spleen cells from a C. trachomatis serovar L2 immunized mouse or a control mouse injected with PBS after a 5 days culture with Cap1#139-147 peptide-coated syngeneic spleen cells and CD8+ Tcells able to specifically recognize Cap1#139-147 epitope gave 73%, 60% and 32% specific lysis at a30:1, 10:1 and 3:1 effector to target ratios, respectively. The control mice had a percent lysis of approximately 10% at a 30:1 effector to target ratio, and steadily declining with lowering E:T ratios. Target cells were Cap1#139-147 peptidecoated, or untreated P815 cells. These data suggest that Cap1#139-147 peptide-specific T-cells are primed during murine infection with C. trachomatis.

Ct529 Localization

Studies were performed demonstrating that Ct529 (referred to herein as

20 Cap-1) localizes to the inclusion membrane of *C. trachomatis*-infected cells and is not
associated with elementary bodies or reticulate bodies. As described above, Cap-1 was
identified as a product from *Chlamydia* that stimulates CD8+ CTL. These CTL are
protective in a murine model of infection, thus making Cap-1 a good vaccine candidate.
Further, since these CTL are MHC-I restricted, the Cap-1 gene must have access to the
25 cytosol of infected cells, which may be a unique characteristic of specific *Chlamydial*gene products. Therefore, determination of the cellular localization of the gene
products would be useful in characterizing Cap-1 as a vaccine candidate. To detect the
intracellular localization of Cap-1, rabbit polyclonal antibodies directed against a
recombinant polypeptide encompassing the N-terminal 125 amino acids of Cap-1 (SEQ
30 ID NO: 305, with the amino acid sequence including the N-terminal 6-His tag provided
in SEO ID NO: 304) were used to stain McCoy cells infected with *Chlamydiae*.

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Rabbit-anti-Cap-1 polyclonal antibodies were obtained by hyperimmunization of rabbits with a recombinant polypeptide, rCt529c1-125 (SEQ ID NO: 305) encompassing the N-terminal portion of Cap-1. Recombinant rCt529c1-125 protein was obtained from *E. coli* transformed with a pET expression plasmid (as described above) encoding the nucleotides 1-375 encoding the N-terminal 1-125 amino acids of Cap-1. Recombinant protein was purified by Ni-NTA using techniques well known in the art. For a positive control antiserum, polyclonal antisera directed against elementary bodies were made by immunization of rabbits with purified *C. trachomatis* elementary bodies (Biodesign, Sacco, Maine). Pre-immune sera derived from rabbits prior to immunization with the Cap-1 polypeptide was used as a negative control.

Immunocytochemistry was performed on McCoy cell monolayers grown on glass coverslips inoculated with either *C. trachomatis* serovar L2 or *C. psitacci*, strain 6BC, at a concentration of 10⁶ IFU (Inclusion Forming Units) per ml. After 2 hours, medium was aspirated and replaced with fresh RP-10 medium supplemented with cycloheximide (1.0 µg/ml). Infected cells were incubated at in 7% CO₂ for 24 hours and fixed by aspirating medium, rinsing cells once with PBS and methanol fixation for 5 minutes. For antigen staining, fixed cell monolayers were washed with PBS and incubated at 37°C for 2 hours with 1:100 dilutions of specific or control antisera. Cells were rinsed with PBS and incubated for 1 hour with fluorescein isothiocyanate (FITC)-labeled, anti-rabbit IgG (KPL, Gaithersburg) and stained with Evans blue (0.05%) in PBS. Fluorescence was observed with a 100X objective (Zeiss epifluorescence microscope), and photographed (Nikon UFX-11A camera).

Results from this study show Cap-1 localizes to the inclusion membrane of *C. trachomatis*-infected cells. Cap-1 specific antibody labeled the inclusion membranes of *C. trachomatis*-infected cells, but not *Chlamydial* elementary bodies contained in these inclusions or released by the fixation process. Conversely, the anti-elementary body antibody clearly labeled the bacterial bodies, not only within the inclusions, but those released by the fixation process. Specificity of the anti-Cap-1 antibody is demonstrated by the fact that it does not stain *C. psittaci-*infected cells. Specificity of the Cap-1 labeling is also shown by the absence of reactivity in pre-immune sera. These results suggest that Cap-1 is released from the bacteria and becomes associated with the *Chlamydial* inclusion membrane. Therefore, Cap-1 is

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gene product which may be useful for stimulating CD8+ T cells in the development of a vaccine against infections caused by *Chlamydia*.

The relevance of the Cap-1 gene as a potential CTL antigen in a vaccine against Chlamydia infection is further illustrated by two additional series of studies.

5 First, CTL specific for the MHC-I epitope of Cap-1 CT529 #138-147 peptide of C. trachomatis (SEQ ID NO: 144) have been shown to be primed to a high frequency during natural infection. Specifically, Balb/C mice were inoculated with 10⁶ LF.U. of C. trachomatis, serova L2. After 2 weeks, spleens were harvested and quantified by Elispot analysis for the number of IFN-\gamma secreting cells in response to Cap-1 #138-147 peptide-pulsed antigen presenting cells. In two experiments, the number of IFN-\gamma secreting cells in 10⁵ splenocytes was about 1% of all CD8+ T-cells. This high frequency of responding CD8+ CTL to the MHC-1 epitope (Cap-1 CT529 #138-147 peptide) suggest that Cap-1 is highly immunogenic in infections.

Results from a second series of studies have shown that the Cap-1 protein is almost immediately accessible to the cytosol of the host cell upon infection. This is shown in a time-course of Cap-1 CT529 #138-147 peptide presentation. Briefly, 3T3 cells were infected with C. trachomatis serovar L2 for various lengths of time, and then tested for recognition by Cap-1 CT529 #138-147 peptide-specific CTL. The results show that C. trachomatis-infected 3T3 cells are targeted for recognition by the antigen-specific CTL after only 2 hours of infection. These results suggest that Cap-1 is an early protein synthesized in the development of C. trachomatis elementary bodies to reticulate bodies. A CD8+ CTL immune response directed against a gene product expressed early in infection may be particularly efficacious in a vaccine against Chlamydia infection.

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EXAMPLE 5

GENERATION OF ANTIBODY AND T-CELL RESPONSES IN MICE IMMUNIZED WITH CHLAMYDIA ANTIGENS

30 Immunogenicity studies were conducted to determine the antibody and CD4+ T cell responses in mice immunized with either purified SWIB or S13 proteins formulated with Montanide adjuvant, or DNA-based immunizations with pcDNA-3 expression

vectors containing the DNA sequences for SWIB or S13. SWIB is also referred to as clone 1-B1-66 (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEO ID NO: 5), and \$13 ribosomal protein is also referred to as clone 10-C10-31 (SEO ID NO: 4, with the corresponding amino acid sequence provided in SEQ ID NO: 12). 5 In the first experiment, groups of three C57BL/6 mice were immunized twice and monitored for antibody and CD4+ T-cell responses. DNA immunizations were intradermal at the base of the tail and polypeptide immunizations were administered by subcutaneous route. Results from standard 3H-incorporation assays of spleen cells from immunized mice shows a strong proliferative response from the group immunized with purified recombinant SWIB polypeptide (SEQ ID NO: 5). Further analysis by cytokine induction assays, as previously described, demonstrated that the group immunized with SWIB polypeptide produced a measurable IFN-y and IL-4 response. Subsequent ELISA-based assays to determine the predominant antibody isotype response in the experimental group immunized with the SWIB polypeptide were performed. Fig. 4 15 illustrates the SWIB-immunized group gave a humoral response that was predominantly IgG1.

In a second experiment, C3H mice were immunized three times with 10 μg purified SWIB protein (also referred to as clone 1-B1-66, SEQ ID NO: 5) formulated in either PBS or Montanide at three week intervals and harvested two weeks after the third immunization. Antibody titers directed against the SWIB protein were determined by standard ELISA-based techniques well known in the art, demonstrating the SWIB protein formulated with Montanide adjuvant induced a strong humoral immune response. T-cell proliferative responses were determined by a XTT-based assay (Scudiero, et al, Cancer Research, 1988, 48:4827). As shown in Fig. 5, splenocytes from mice immunized with the SWIB polypeptide plus Montanide elicited an antigen specific proliferative response. In addition, the capacity of splenocytes from immunized animals to secrete IFN-γ in response to soluble recombinant SWIB polypeptide was determined using the cytokine induction assay previously described. The splenocytes from all animals in the group immunized with SWIB polypeptide formulated with montanide adjuvant secreted IFN-γ in response to exposure to the SWIB Chlamydia antigen, demonstrating an Chlamydia-specific immune response.

In a further experiment, C3H mice were immunized at three separate time points at the base of the tail with 10 μg of purified SWIB or S13 protein (C. trachomatis, SWIB protein, clone 1-B1-66, SEQ ID NO: 5, and S13 protein, clone 10-C10-31, SEQ ID NO: 4) formulated with the SBAS2 adjuvant (SmithKline Beecham, London, England). Antigen-specific antibody titers were measured by ELISA, showing both polypeptides induced a strong IgG response, ranging in titers from 1 x10⁻⁴ to 1 x10⁻⁵. The IgG1 and IgG2a components of this response were present in fairly equal amounts. Antigen-specific T-cell proliferative responses, determined by standard ³H-incorporation assays on spleen cells isolated from immunized mice, were quite strong for SWIB (50,000 cpm above the negative control) and even stronger for s13 (100,000 cpm above the negative control). The IFNγ production was assayed by standard ELISA techniques from supernatant from the proliferating culture. In vitro restimulation of the culture with S13 protein induced high levels of IFNγ production, approximately 25 ng/ml versus 2 ng/ml for the negative control. Restimulation with the SWIB protein also induced IFNγ, although to a lesser extent.

In a related experiment, C3H mice were immunized at three separate time points with 10 µg of purified SWIB or S13 protein (*C. trachomatis*, SWIB protein, clone 1-B1-66, SEQ ID NO: 5, and S13 protein, clone 10-C10-31, SEQ ID NO: 4) mixed with 10 µg of Cholera Toxin. Mucosal immunization was through intranasal inoculation. Antigen-specific antibody responses were determined by standard ELISA techniques. Antigen-specific IgG antibodies were present in the blood of SWIB-immunized mice, with titers ranging from 1 x10⁻³ to 1 x10⁻⁴, but non-detectable in the S13-immunized animals. Antigen-specific T-cell responses from isolated splenocytes, as measured by IFNy production, gave similar results to those described immediately above for systemic immunization.

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An animal study was conducted to determine the immunogenicity of the
CT529 serovar LGVII CTL epitope, defined by the CT529 10mer consensus peptide
(CSFIGGITYL – SEQ ID NO: 31), which was identified as an H2-Kd restricted CTL
epitope. BALB/c mice (3 mice per group) were immunized three times with 25 µg of
peptide combined with various adjuvants. The peptide was administered systemically at
the base of the tail in either SKB Adjuvant System SBAS-2". SBAS-7 (SmithKline

Beecham, London, England) or Montanide. The peptide was also administered intranasally mixed with 10ug of Cholera Toxin (CT). Naive mice were used as a control. Four weeks after the 3rd immunization, spleen cells were restimulated with LPS-blasts pulsed with 10ug/ml CT529 10mer consensus peptide at three different effector to LPS-blasts ratios: 6, 1.5 and 0.4 at 1x10⁶ cell/ml. After 2 restimulations, effector cells were tested for their ability to lyse peptide pulsed P815 cells using a standard chromium release assay. A non-relevant peptide from chicken egg ovalbumin was used as a negative control. The results demonstrate that a significant immune response was elicited towards the CT529 10mer consensus peptide and that antigen-specific T-cells capable of lysing peptide-pulsed targets were elicited in response to immunization with the peptide. Specifically, antigen-specific lytic activities were found in the SBAS-7 and CT adjuvanted group while Montanide and SBAS-2" failed to adjuvant the CTL epitope immunization.

15 EXAMPLE 6

EXPRESSION AND CHARACTERIZATION OF CHLAMYDIA PNEUMONIAE GENES

The human T-cell line, TCL-8, described in Example 1, recognizes

20 Chiamydia trachomatis as well as Chiamydia pneumonia infected monocyte-derived dendritic cells, suggesting Chiamydia trachomatis and pneumonia may encode cross-reactive T-cell epitopes. To isolate the Chiamydia pneumonia genes homologous to Chiamydia trachomatis LGV II clones 1B1-66, also referred to as SWIB (SEQ ID NO: 1) and clone 10C10-31, also referred to as S13 ribosomal protein (SEQ ID NO: 4),

25 HeLa 229 cells were infected with C. pneumonia strain TWAR (CDC/CWL-029). After three days incubation, the C. pneumonia-infected HeLa cells were harvested, washed and resuspended in 200 µl water and heated in a boiling water bath for 20 minutes. Ten microliters of the disrupted cell suspension was used as the PCR template.

C. pneumonia specific primers were designed for clones 1B1-66 and 10C10-31 such that the 5' end had a 6X-Histidine tag and a Nde I site inserted, and the 3' end had a stop codon and a BamHI site included (Fig. 6). The PCR products were amplified and sequenced by standard techniques well known in the art. The C.

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pneumonia-specific PCR products were cloned into expression vector pET17B (Novagen, Madison, WI) and transfected into E. coli BL21 pLysS for expression and subsequent purification utilizing the histidine-nickel chromatographic methodology provided by Novagen. Two proteins from C. pneumonia were thus generated, a 10-11 kDa protein referred to as CpSWIB (SEQ ID NO: 27, and SEQ ID NO: 78 having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 77, having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 30 and 91, respectively).

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EXAMPLE 7

INDUCTION OF T CELL PROLIFERATION AND INTERFERON-Y PRODUCTION BY CHLAMYDIA PNEUMONIAE ANTIGENS

The ability of recombinant *Chlamydia pneumoniae* antigens to induce T cell proliferation and interferon-y production is determined as follows.

Proteins are induced by IPTG and purified by Ni-NTA agarose affinity chromatography (Webb et al., *J. Immunology 157:*5034-5041, 1996). The purified polypeptides are then screened for the ability to induce T-cell proliferation in PBMC 20 preparations. PBMCs from *C. pneumoniae* patients as well as from normal donors whose T-cells are known to proliferate in response to *Chlamydia* antigens, are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 µg/ml gentamicin. Purified polypeptides are added in duplicate at concentrations of 0.5 to 10 µg/mL. After six days of culture in 96-well round-bottom plates in a volume 25 of 200 µl, 50 µl of medium is removed from each well for determination of IFN-γ levels, as described below. The plates are then pulsed with 1 µCi/well of tritiated thymidine for a further 18 hours, harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

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IFN-y was measured using an enzyme-linked immunosorbent assay (ELISA). ELISA plates are coated with a mouse monoclonal antibody directed to human IFN-y (PharMingen, San Diego, CA) in PBS for four hours at room temperature. Wells are then blocked with PBS containing 5% (W/V) non-fat dried milk for 1 hour at room temperature. The plates are washed six times in PBS/0.2% TWEEN-20 and samples diluted 1:2 in culture medium in the ELISA plates are incubated overnight at room temperature. The plates are again washed and a polyclonal rabbit anti-human IFN-y serum diluted 1:3000 in PBS/10% normal goat serum is added to each well. The plates are then incubated for two hours at room temperature, washed and horseradish peroxidase-coupled anti-rabbit IgG (Sigma Chemical So., St. Louis, MO) is added at a 1:2000 dilution in PBS/5% non-fat dried milk. After a further two hour incubation at room temperature, the plates are washed and TMB substrate added. The reaction is stopped after 20 min with 1 N sulfuric acid. Optical density is determined at 450 nm using 570 nm as a reference wavelength. Fractions that result in both replicates giving an OD two fold greater than the mean OD from cells cultured in medium alone, plus 3 standard deviations, are considered positive.

A human anti-Chlamydia T-cell line (TCL-8) capable of cross-reacting to C. trachomatis and C. pneumonia was used to determine whether the expressed proteins described in the example above, (i.e., CpSWIB, SEO ID NO; 27, and SEO ID NO: 78 having a 6X His tag, with the corresponding amino acid sequence provided in SEQ ID NO: 28, respectively, and the 15 kDa protein referred to as CpS13 SEQ ID NO: 29, and SEO ID NO: 77, having a 6X His tag, with the corresponding amino acid sequence provided in SEO ID NO: 30 and 91, respectively), possessed T-cell epitopes common to both C. trachomatis and C. pneumonia. Briefly, E. coli expressing 25 Chlamydial proteins were titered on 1 x 10⁴ monocyte-derived dendritic cells. After two hours, the dendritic cells cultures were washed and 2.5 x 104 T cells (TCL-8) added and allowed to incubate for an additional 72 hours. The amount of INF-y in the culture supernatant was then determined by ELISA. As shown in Figs. 7A and 7B, the TCL-8 T-cell line specifically recognized the S13 ribosomal protein from both C. trachomatis and C. pneumonia as demonstrated by the antigen-specific induction of IFN-y, whereas 30 only the SWIB protein from C. trachomatis was recognized by the T-cell line. To

validate these results, the T cell epitope of C, trachomatis SWIB was identified by epitone mapping using target cells pulsed with a series of overlapping pertides and the T-cell line TCL-8. 3H-thymidine incorporation assays demonstrated that the peptide, referred to as C.t.SWIB 52-67, of SEO ID NO: 39 gave the strongest proliferation of the 5 TCL-8 line. The homologous peptides corresponding to the SWIB of C. pneumoniae sequence (SEQ ID NO: 40), the topoisomerase-SWIB fusion of C. pneumoniae (SEQ ID NO: 43) and C. trachomatis (SEO ID NO: 42) as well as the human SWI domain (SEQ ID NO: 41) were synthesized and tested in the above assay. The T-cell line TCL-8 only recognized the C. trachomatis peptide of SEQ ID NO: 39 and not the 10 corresponding C. pneumoniae peptide (SEO ID NO: 40), or the other corresponding peptides described above (SEQ ID NO; 41-43).

Chlamydia-specific T cell lines were generated from donor CP-21 with a positive serum titer against C. pneumoniae by stimulating donor PBMC with either C. trachomatis or C. pneumoniae-infected monocyte-derived dendritic cells, respectively. 15 T-cells generated against C. pneumoniae responded to recombinant C. pneumoniae-SWIB but not C. trachomatis-SWIB, whereas the T-cell line generated against C. trachomatis did not respond to either C. trachomatis- or C. pneumoniae-SWIB (see Fig. 9). The C. pneumoniae-SWIB specific immune response of donor CP-21 confirms the C. pneumoniae infection and indicates the elicitation of C. pneumoniae-SWIB specific T-cells during in vivo C. pneumoniae infection.

Epitope mapping of the T-cell response to C. pneumoniae-SWIB has shown that Cp-SWIB-specific T-cells responded to the overlapping peptides Cp-SWIB 32-51 (SEO ID NO: 101) and Cp-SWIB 37-56 (SEO ID NO: 102), indicating a C. pneumoniae-SWIB-specific T-cell epitope Cp-SWIB 37-51 (SEQ ID NO: 100).

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In additional experiments, T-cell lines were generated from donor CP1, also a C. pneumoniae seropositive donor, by stimulating PBMC with non-infectious elementary bodies from C. trachomatis and C. pneumoniae, respectively. In particular, proliferative responses were determined by stimulating 2.5 x 104 T-cells in the presence of 1 x 104 monocyte-derived dendritic cells and non-infectious elementary bodies 30 derived from C. trachomatis and C. pneumoniae, or either recombinant C. trachomatis or C. pneumoniae SWIB protein. The T-cell response against SWIB resembled the data obtained with T-cell lines from CP-21 in that C. pneumoniae-SWIB, but not C.

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trachomatis-SWIB elicited a response by the C. pneumoniae T-cell line. In addition, the C. trachomatis T-cell line did not proliferate in response to either C. trachomatis or C. pneumoniae SWIB, though it did proliferate in response to both CT and CP elementary bodies. As described in Example 1, Clone 11-C12-91 (SEQ ID NO: 63), 5 identified using the TCP-21 cell line, has a 269 bp insert that is part of the OMP2 gene (CT443) and shares homology with the 60 kDa cysteine rich outer membrane protein of C. pneumoniae, referred to as OMCB. To further define the reactive epitope(s), epitope mapping was performed using a series of overlapping peptides and the immunoassay previously described. Briefly, proliferative responses were determined by stimulating 10 2.5 x 10⁴ TCP-21 T-cells in the presence of 1 x 10⁴ monocyte-derived dendritic cells with either non-infectious elementary bodies derived from C. trachomatis and C. pneumoniae, or peptides derived from the protein sequence of C. trachomatis or C. pneumoniae OMCB protein (0.1 µg/ml). The TCP-21 T-cells responded to epitopes CT-OMCB #167-186, CT-OMCB #171-190, CT-OMCB #171-186, and to a lesser 15 extent, CT-OMCB #175-186 (SEQ ID NO: 249-252, respectively). Notably, the TCP-21 T-cell line also gave a proliferative response to the homologous C. pneumoniae peptide CP-OMCB #171-186 (SEO ID NO: 253), which was equal to or greater than the response to the to the C. trachomatis peptides. The amino acid substitutions in position two (i.e., Asp for Glu) and position four (i.e., Cys for Ser) did not alter the proliferative 20 response of the T-cells and therefore demonstrating this epitope to be a cross-reactive epitope between C. trachomatis and C. pneumoniae.

EXAMPLE 8

IMMUNE RESPONSES OF HUMAN PBMC AND T-CELL LINES AGAINST CHLAMYDIA ANTIGENS

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The examples provided herein suggest that there is a population of healthy donors among the general population that have been infected with C. trachomatis and generated a protective immune response controlling the C. trachomatis infection. These donors remained clinically asymptomatic and seronegative for C. trachomatis. To characterize the immune responses of normal donors against chiamydial antigens which had been identified by CD4 expression cloning, PBMC

obtained from 12 healthy donors were tested against a panel of recombinant chlamydial antigens including C. trachomatis-, C. pneumoniae-SWIB and C. trachomatis-, C. pneumoniae-SI3. The data are summarized in Table I below. All donors were seronegative for C. trachomatis, whereas 6/12 had a positive C. pneumoniae titer.

5 Using a stimulation index of >4 as a positive response, 11/12 of the subjects responded to C. trachomatis elementary bodies and 12/12 responded to C. pneumoniae elementary bodies. One donor, AD104, responded to recombinant C. pneumoniae-SI3 protein, but not to recombinant C. trachomatis-SI3 protein, indicating a C. pneumoniae-specific response. Three out of 12 donors had a C. trachomatis-SWIB, but not a C. pneumoniae-SWIB specific response, confirming a C. trachomatis infection. C. trachomatis and C. pneumoniae-SI3 elicited a response in 8/12 donors suggesting a chlamydial infection. These data demonstrate the ability of SWIB and SI3 to elicit a T-cell response in PBMC of normal study subjects.

120 Table I.

Immune response of normal study subjects against Chlamalia

Donor	Sex	<i>Chlamydia</i> IgGtiter	CT EB	CP EB	CT Swib	CP Swib	CT Sl3	CP SI3	CT lpdA	CT TSA
4D100	male	negative	++	+++	+		++	++		n.t.
4D104	female	negative	+++	++	_	_	_	++		nt.
4D108	male	CP 1:256	++	++	+	+/-	+	+	+	nt.
4D112	ferrale	negative	++	++	+	-	+		+/-	nt
AD120	male	negative	-	+	-	-	-	-	-	nt.
4D124	female	CP 1:128	++	++	-	-	-	-	-	nt
4D128	male	CP 1:512	+	++	-	-	++	+	++	-
4D132	female	negative	++	++	-	-	+	+	-	-
4D136	female	CP 1:128	+	++	-	-	+/-	-	-	-
4D140	male	CP 1:256	++	++	-	-	+	+	-	-
4D142	female	CP 1:512	++	++	-	-	+	+	+	-
4D146	fermle	negative	++	++	_	-	++	+	+	_

CT= Chlamydia trachomatis; CP= Chlamydia pneumoniae; EB= Chlamydia elementary bodies; Swib= recombinant Chlamydia Swib protein; S13= recombinant Chlamydia S13 protein; pdA= recombinant Chlamydia pldA protein; TSA= recombinant Chlamydia TSA protein. Values represent results from standard proliferation assays. Proliferative responses were determined by stimulating 3 x 10⁵ pBMC with 1 x 10⁴ monocyte-derived dendritic cells pre-incubated with the respective recombinant antigens or elementary bodies (EB). Assays were harvested after 6 days with a ³H-thymidine pulse for the last 18h.

SI: Stimulation index

+/-: SI ~ 4 15 +: SI > 4 ++: SI 10-30 +++: SI 30

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In a first series of experiments, T-cell lines were generated from a healthy female individual (CT-10) with a history of genital exposure to C. trachomatis by stimulating T-cells with C. trachomatis LGV II elementary bodies as previously described. Although the study subject was exposed to C. trachomatis, she did not seroconvert and did not develop clinical symptoms, suggesting donor CT-10 may have developed a protective immune response against C. trachomatis. As shown in Fig. 10, a primary Chlamydia-specific T-cell line derived from donor CT-10 responded to C. trachomatis-SWIB, but not C. pneumoniae-SWIB recombinant proteins, confirming the exposure of CT-10 to C. trachomatis. Epitope mapping of the T-cell response to C. trachomatis-SWIB showed that this donor responded to the same epitope Ct-SWIB 52-67 (SEO ID NO: 39) as T-cell line TCL-8, as shown in Fig. 11.

Additional T-cell lines were generated as described above for various C. trachomatis patients. A summary of the patients' clinical profile and proliferative responses to various C. trachomatis and C. pneumoniae elementary bodies and recombinant proteins are summarized in Table II as follows:

	Prolifer	rative respon	nse of	C. tre	achom	<i>atis</i> pa	tient	s		
Patients	Clinical manifestation	IgG titer	CT EB	CP EB	CT Swib	CP Swib	CT S13	CP S13	CT lpdA	CT TSA
CT-1	NGU	negative	+	+	-	-	++	++	++	+
CT-2	NGU	negative	++	++	-	-	+	+/-	-	-
СТ-3	asymptomatic shed Eb Dx was HPV	Ct 1:512 Cp 1:1024 Cps 1:256	+	+	-	-	+	-	+	-
CT-4	asymptomatic shed Eb	Ct 1:1024	+	+	-	-	-	-	-	-
CT-5	BV	Ct 1:256 Cp 1:256	++	++	-	-	+	-	-	-
CT-6	perinial rash discharge	Cp 1:1024	+	+	-	-	-	-	-	-
CT-7	BV genital ulcer	Ct 1:512 Cp 1:1024	+	+	-	-	+	+	+	-
CT-8	Not known	Not tested	++	++	-	-	-	-	-	-
CT-9	asymptomatic	Ct 1:128 Cp 1:128	+++	++	-	-	++	+	+	-
CT-10	Itch mild vulvar	negative	++	++	-	-	-	-	-	-
CT-11	BV, abnormal pap	Ct 1: 512	+++	+++	-	-	+++	+/-	++	+
CT-12	asymptomatic	Cp 1: 512	++	++	-	-	++	+	+	_

NGU= Non-Gonococcal Urethritis; BV= Bacterial Vaginosis; CT= Chlamydia trachomatis; CP= Chlamydia pneumoniae; EB= Chlamydia elementary bodies; Swib= recombinant Chlamydia Swib protein; S13= recombinant Chlamydia S13 protein; lpdA= recombinant Chlamydia IpdA protein; TSA= recombinant Chlamydia TSA protein

Values represent results from standard proliferation assays. Proliferative responses were determined by stimulating 3 x 10³ PBMC with the respective recombinant antigens or elementary bodies (EB). Assays were harvested after 6 days with a ³H-thymidine pulse for the last 18 hours.

SI: Stimulation index

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Using the panel of asymptomatic (as defined above) study subjects and C. trachomatis patients, as summarized in Tables I and II, a comprehensive study of the immune responses of PBMC derived from the two groups was conducted. Briefly, PBMCs from C. pneumoniae patients as well as from normal donors are cultured in medium comprising RPMI 1640 supplemented with 10% pooled human serum and 50 u g/ml gentamicin. Purified polypeptides, a panel of recombinant chlamydial antigens including C. trachomatis-, C. pneumoniae-SWIB and S13, as well as , C. trachomatis lpdA and TSA are added in duplicate at concentrations of 0.5 to 10 ug/mL. After six days of culture in 96-well round-bottom plates in a volume of 200 µl, 50 µl of medjum is removed from each well for determination of IFN-y levels, as described below. The plates are then pulsed with 1 uCi/well of tritiated thymidine for a further 18 hours. harvested and tritium uptake determined using a gas scintillation counter. Fractions that result in proliferation in both replicates three fold greater than the proliferation observed in cells cultured in medium alone are considered positive.

Proliferative responses to the recombinant Chlamydiae antigens demonstrated that the majority of asymptomatic donors and C. trachomatis patients recognized the C. trachomatis S13 antigen (8/12) and a majority of the C. trachomatis patients recognized the C. pneumonia S13 antigen (8/12), with 4/12 asymptomatic donors also recognizing the C. pneumonia S13 antigen. Also, six out of twelve of the C. trachomatis patients and four out of twelve of the asymptomatic donors gave a proliferative response to the lpdA antigen of C. trachomatis. These results demonstrate that the C. trachomatis and C. pneumonia S13 antigen, C. trachomatis Swib antigen and the C. trachomatis lpdA antigen are recognized by the asymptomatic donors, indicating these antigens were recognized during exposure to Chlamvdia and an immune response elicited against them. This implies these antigens may play a role in conferring 25 protective immunity in a human host. In addition, the C. trachomatis and C. pneumonia S13 antigen is recognized equally well among the C. trachomatis patients. therefore indicating there may be epitopes shared between C. trachomatis and C. pneumonia in the S13 protein. Table III summarizes the results of these studies.

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Table III.

Antigen	Normal Donors	C.t. Patients
C.tSwib	3/12	0/12
C.pSwib	0/12	0/12
C.tS13	8/12	8/12
C.pS13	4/12	8/12
lpdA	4/12	6/12
TSA	0/12	2/12

A series of studies were initiated to determine the cellular immune response to short-term T-cell lines generated from asymptomatic donors and C. trachomatis patients. Cellular immune responses were measured by standard proliferation assays and IFN-y, as described in Example 7. Specifically, the majority of the antigens were in the form of single E. coli clones expressing Chlamydial antigens, although some recombinant proteins were also used in the assays. The single E. coli clones were titered on 1 x 104 monocyte-derived dendritic cells and after two hours, the culture was washed and 2.5 x 104 T-cells were added. The assay using the recombinant proteins were performed as previously described. Proliferation was determined after four days with a standard 3H-thymidine pulse for the last 18 hours. Induction of IFN-y 15 was determined from culture supernatants harvested after four days using standard ELISA assays, as described above. The results show that all the C. trachomatis antigens tested, except for C.T. Swib, elicited a proliferative response from one or more different T-cell lines derived form C. trachomatis patients. In addition, proliferative responses were elicited from both the C. trachomatis patients and asymptomatic donors for the following Chlamydia genes, CT622, groEL, pmpD, CT610 and rS13.

The 12G3-83 clone also contains sequences to CT734 and CT764 in addition to CT622, and therefore these gene sequence may also have immunoreactive epitopes. Similarly, clone 21G12-60 contains sequences to the hypothetical protein genes CT229 and CT228 in addition to CT875; and 15H2-76 also contains sequences from CT812 and CT088, as well as sharing homology to the sycE gene. Clone 11H3-61 also contains sequences sharing homology to the PGP6-D virulence protein.

Table IV.

Clone	C. t.	TCL from	TCL from	SEQ ID NO:
	Antigen	Asymp. Donors	C. t.	
	(putative*)		Patients	
1B1-66 (E. coli)	Swib	2/2	0/4	5
1B1-66 (protein)	Swib	2/2	0/4	5
12G3-83 (E. coli)	CT622*	2/2	4/4	57
22B3-53 (E. coli)	· groEL	1/2	4/4	111
22B3-53 (protein)	groEL	1/2	4/4	111
15H2-76 (E. coli)	PmpD*	1/2	3/4	87
11H3-61 (E. coli)	rL1*	0/2	3/4	60
14H1-4 (E. coli)	TSA	0/2	3/4	56
14H1-4 (protein)	TSA	0/2	3/4	56
11G10-46 (E. coli)	CT610	1/2	1/4	62
10C10-17 (E. coli)	rS13	1/2	1/4	62
10C10-17 (protein)	rS13	1/2	1/4	62
21G12-60 (E. coli)	CT875*	0/2	2/4	110
11H4-32 (E. coli)	dnaK	0/2	2/4	59
21C7-8 (E. coli)	dnaK	0/2	2/4	115
17C10-31 (E. coli)	CT858	0/2	2/4	114

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EXAMPLE 9

PROTECTION STUDIES USING CHLAMYDIA ANTIGENS

1. SWIB

Protection studies were conducted in mice to determine whether immunization with chlamydial antigens can impact on the genital tract disease resulting from chlamydial inoculation. Two models were utilized; a model of intravaginal inoculation

that uses a human isolate containing a strain of Chlamydia psittaci (MTW447), and a model of intrauterine inoculation that involves a human isolate identified as Chlamvdia trachomatis, serovar F (strain NI1). Both strains induce inflammation in the upper genital tract, which resemble endometritis and salpingitis caused by Chlamydia 5 trachomatis in women. In the first experiment, C3H mice (4 mice per group) were immunized three times with 100 ug of pcDNA-3 expression vector containing C. trachomatis SWIB DNA (SEO ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5). Inoculations were at the base of the tail for systemic immunization. Two weeks after the last immunization, animals were progesterone treated and infected, either thru the vagina or by injection of the inoculum in the uterus. Two weeks after infection, the mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. Inflammation level was scored (from + for very mild, to +++++ for very severe). Scores attributed to each single oviduct/ovary were summed and divided by the number of organs examined to get a mean score of 15 inflammation for the group. In the model of uterine inoculation, negative controlimmunized animals receiving empty vector showed consistent inflammation with an ovary/oviduct mean inflammation score of 6.12, in contrast to 2.62 for the DNAimmunized group. In the model of vaginal inoculation and ascending infection, negative control-immunized mice had an ovary /oviduct mean inflammation score of 8.37, versus 5.00 for the DNA-immunized group. Also, in the later model, vaccinated mice showed no signs of tubal occlusion while negative control vaccinated groups had inflammatory cells in the lumen of the oviduct

In a second experiment, C3H mice (4 mice per group) were immunized three times with 50 µg of pcDNA-3 expression vector containing *C. trachomatis* SWIB

25 DNA (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEQ ID NO: 5) encapsulated in Poly Lactide co-Glycolide microspheres (PLG); immunizations were made intra-peritoneally. Two weeks after the last immunization, animal were progesterone treated and infected by inoculation of *C. psittaci* in the vagina. Two weeks after infection, mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. Inflammation level was scored as previously described. Scores attributed to each single oviduct lovary were summed and divided by the number of examined organs to get a mean of inflammation for the group. Negative control-

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immunized animals receiving PLG-encapsulated empty vector showed consistent infammation with an ovary /oviduct mean inflammation score of 7.28, versus 5.71 for the PLG-encapsulated DNA immunized group. Inflammation in the peritoneum was 1.75 for the vaccinated group versus 3.75 for the control.

In a third experiment, C3H mice (4 per group) were immunized three times with 10 µg of purified recombinant protein, either SWIB (SEQ ID NO: 1, with the corresponding amino acid sequence provided in SEO ID NO: 5, or S13 (SEO ID NO: 4, with the corresponding amino acid sequence provided in SEQ ID NO: 12) mixed with Cholera Toxin (CT); the preparation was administred intranasally upon anaesthesia in a 20 uL volume. Two weeks after the last immunization, animal were progesterone treated and infected, either by vaginal inoculation of C. psittaci or by injection of C. trachomatis serovar F in the uterus. Two weeks after infection, the mice were sacrificed and genital tracts sectioned, stained and examined for histopathology. The degree of inflammation was scored as described above. Scores attributed to each single oviduct/ovary were summed and divided by the number of examined organs to get a mean score of inflammation for the group. In the model of uterine inoculation, negative control- immunized animals receiving cholera toxin alone showed an ovary/oviduct mean inflammation score of 4.25 (only 2 mice analyzed; 2 other died) versus 5.00 for the s13 plus cholera toxin-immunized group, and 1.00 for the SWIB plus cholera toxin. Untreated infected animals had an ovary /oviduct mean inflammation score of 7. In the model of vaginal inoculation and ascending infection, negative control-immunized mice had an ovary /oviduct mean inflammation score of 7.37 versus 6.75 for the s13 plus cholera toxin-immunized group and 5.37 for the SWIB plus cholera toxin-immunized group. Untreated infected animals had an ovary /oviduct mean inflammation score of 8.

The three experiments described above suggest that SWIB-specific protection is obtainable. This protective effect is more marked in the model of homologous infection but is still present when in a heterologous challenge infection with C. psittaci.

CT529/Cap1

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CT529/Cap1 was identified earlier as a product from Chlamydia that stimulates CD8+ CTL. In this example, we sought to confirm that immunization with Cap1 would be protective in an animal model of chlamydia infection.

To generate recombinant vaccinia virus for delivery of a Cap1 immunogenic fragment, a DNA fragment containing a modified Kozak sequence and base pairs 319-530 of the cap1 gene (CT529) was amplified from C. trachomatis L2 genomic DNA using PCRTM and ligated into pSC11ss (Earl PL, Koenig S, Moss B (1991) Biological and immunological properties of human immunodeficiency virus type 1 envelope 10 glycoprotein: analysis of proteins with truncations and deletions expressed by recombinant vaccinia viruses. J Virol. 65:31-41). DNA digested with Sall and Stul. The portion of the cap1 gene ligated into pSC11ss encodes amino acids 107-176 of Cap1 protein, containing the previously identified CTL epitope of amino acids 139-147. The resulting plasmid was used to transfect CV-1 cells (ATCC# CCL-70; Jensen FC et al. (1964) Infection of human and simian tissue cultures with Rous Sarcoma Virus. Proc. Natl. Acad. Sci. USA 52: 53-59.) which were subsequently infected with wildtype vaccinia virus. Homologous recombination between the wild-type virus and plasmid DNA generated recombinant vaccinia viruses which were selected on the basis of both beta-galactosidase expression and the inactivation of thymidine kinase, as 20 described previously (Chakrabarti et al, Mol Cell Biol. 1985, 5(12):3403-9). Recombinant virus was plaque purified three times and titered after growth in human TK-143B cells. Virus preparations were treated with equal volume of 0.25 mg/ml trypsin for 30 mins, at 37°C and diluted in PBS prior to immunization of mice. Groups of 5 mice were used for all experimental and control groups. The data presented below are representative of three independent experiments.

A group of mice was immunized with 106 of the recombinant vaccinia i.p. and was allowed to recover for 3 weeks. Negative control groups were immunized with either buffer alone or wild-type vaccinia. As a positive control, a group of mice was infected i.v. with 106 i.f.u. of C. trachomatis. The number of organisms given to the positive control group has been previously shown to be cleared within 2 weeks. After 3 weeks, animals in each of the groups were challenged i.v. with 106 i.f.u. of C. trachomatis. Three days after challenge the mice were sacrificed and the number of i.f.u. per spleen was determined.

The mean number of organisms found in the spleens of animals immunized with
the vaccinia virus expressing Capl (7.1x10⁴) was 2.6-fold fewer (p<0.01; Wilcoxon'sRank Sum analysis) than animals in the control groups immunized with either buffer
(1.8x10⁵) or wild-type vaccinia (1.9x10⁵). Animals in the positive group had 77-fold
fewer organisms (2.4x103) per spleen than animals in the negative control groups
(p<0.01; Wilcoxon's-Rank Sum analysis). These data demonstrate that immunization
with an immunogenic fragment of Capl can afford a statistically significant level of

EXAMPLE 10 Pmp/Ra12 FUSION PROTEINS

Various Pmp/Ra12 fusion constructs were generated by first synthesizing

PCR fragments of a Pmp gene using primers containing a Not I restriction site. Each

PCR fragment was then ligated into the Notl restriction site of pCRX1. The pCRX1

vector contains the 6HisRa12 portion of the fusion. The Ra12 portion of the fusion

construct encodes a polypeptide corresponding to amino acid residues 192-323 of

Mycobacterium tuberculosis MTB32A, as described in U.S. Patent Application

60/158,585, the disclosure of which is incorporated herein by reference. The correct

orientation of each insert was determined by its restriction enzyme pattern and its

sequence was verified. Multiple fusion constructs were made for PmpA, PmpB, PmpC,

PmpF and PmpH, as described further below:

PmpA Fusion Proteins

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PmpA is 107 kD protein containing 982 as and was cloned from serovar E. The PmpA protein was divided into 2 overlapping fragments, the PmpA(N-terminal) and (C-terminal) portions.

PmpA(N-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGTTTATAACAAAGGAACTTATG (SEO ID NO: 306)

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GAGAGCGGCCGCTTACTTAGGTGAGAAGAAGGGAGTTTC (SEQ ID NO: 307)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 308, encoding a 66 kD protein (619aa) expressing the segment 1-473 aa of PmpA. The amino acid sequence of the fusion protein is set forth in SEO ID NO: 309.

PmpA(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCCATTCTATTCATTTCTTTGATCCTG (SEQ ID NO: 310) GAGAGCGGCCGCTTAGAAGCCAACATAGCCTCC (SEQ ID NO: 311)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID

NO: 312, encoding a 74 kD protein (691aa) expressing the segment 438-982 aa of

PmpA. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 313.

PmpF Fusion Proteins

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PmpF is 112 kD protein containing 1034 aa and was cloned from the serovar E. PmpF protein was divided into 2 overlapping fragments, the PmpF(N- term) and (C-term) portions.

PmpF(N-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGATTAAAAGAACTTCTCTATCC (SEQ ID NO: 314)

GAGAGCGGCCGCTTATAATTCTGCATCATCTTCTATGGC (SEQ ID NO: 315)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 316, encoding a 69 kD protein (646aa) expressing the segment 1-499 aa of PmpF. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 317.

PmpF(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGACATACGAACTCTGATGGG (SEQ ID NO: 318)

GAGAGCGGCCGCTTAAAAGACCAGAGCTCCTCC (SEQ ID NO: 319)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 320,
25 encoding a 77 kD protein (715aa) expressing the segment 466-1034aa of PmpF. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 321.

PmpH Fusion Proteins

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PmpH is 108 kD protein containing 1016 aa and was cloned from the serovar E. PmpH protein was divided into 2 overlapping fragments, the PmpH(Nterm)and (C-term)portions.

PmpH(N-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCATGCCTTTTTCTTTGAGATCTAC (SEO ID NO: 322) GAGAGCGCCGCTTACACAGATCCATTACCGGACTG (SEQ ID NO: 323)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 324, encoding a 64 kD protein (631aa) expressing the segment 1-484 aa of PmpH. The amino acid sequence of the fusion protein is set forth in SEO ID NO: 325. The donor line CHH037 was found to be reactive against this protein.

PmpH(C-term) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGATCCTGTAGTACAAAATAATTCAGC (SEQ ID NO: 326) GAGAGCGGCCGCTTAAAAGATTCTATTCAAGCC (SEQ ID NO: 327)

15 respectively. The resulting fusion construct has a DNA sequence set forth in SEO ID NO: 328, encoding a 77 kD protein (715aa) expressing the segment 449-1016aa of PmpH. The amino acid sequence of the fusion protein is set forth in SEO ID NO: 329. The patient line CT12 was found to be reactive in response to this protein.

PmpB Fusion Proteins

PmpB is 183 kD protein containing 1750 aa and was cloned from the serovar E. PmpB protein was divided into 4 overlapping fragments, PmpB(1), (2), (3) and (4).

PmpB(1) was amplified by the sense and antisense primers:

GAGAGCGCCCCTCATGAAATGGCTGTCAGCTACTGCG (SEO ID NO: 330)

25 GAGAGCGGCCGCTTACTTAATGCGAATTTCTTCAAG (SEO ID NO: 331)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 332, and encodes is a 53 kD protein (518aa) expressing the segment 1-372 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 333.

PmpB(2) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGGTGACCTCTCAATTCAATCTTC (SEQ ID NO: 334)

GAGAGCGGCCGCTTAGTTCTCTGTTACAGATAAGGAGAC (SEQ ID NO: 335)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 336 and 5 encodes a 60 kD protein (585aa) expressing the segment 330-767 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 337. Cell lines derived from patient lines CT1, CT3, CT4 responded to this recombinant pmpB protein.

PmpB(3) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGACCAACTGAATATCTCTGAGAAC (SEQ ID NO: 338)

0 GAGCGGCCGCTTAAGAGACTACGTGGAGTTCTG (SEQ ID NO: 339)

respectively. The resulting fusion has a DNA sequence set forth in SEQ ID NO: 340 encodes a 67 kD protein (654aa) expressing the segment 732-1236 aa of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 341

PmpB(4) was amplified by the sense and antisense primers:

15 GAGAGCGGCCGCTCGGAACTATTGTGTTCTCTTCTG (SEQ ID NO: 342)

GAGAGCGCCGCTTAGAAGATCATGCGAGCACCGC (SEQ ID NO: 343)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 344 encodes a 76 kD protein (700aa) expressing the segment 1160-1750 of PmpB. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 345.

20 PmpC Fusion Proteins

PmpC is 187 kD protein containing 1774 aa and was cloned from the serovar E/L2. PmpC protein was divided into 3 overlapping fragments, PmpC(1), (2) and (3).

PmpC(1) was amplified by the sense and antisense primers:

25 GAGAGCGGCCGCTCATGAAATTTATGTCAGCTACTGC (SEQ ID NO: 346)
GAGAGCGGCCGCTTACCCTGTAATTCCAGTGATGGTC (SEQ ID NO: 347)

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respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 348 and encodes a 51 kD protein (487aa) expressing the segment 1-340 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 349.

PmpC(2) was amplified by the sense and antisense primers:

GAGAGCGGCCGCTCGATACACAAGTATCAGAATCACC (SEQ ID NO: 350)
GAGAGCGGCCGCTTAAGAGGACGATGAGACACTCTCG (SEO ID NO: 351)

respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID NO: 352 and encodes a 60 kD protein (583aa) expressing the segment 305-741 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEQ ID NO: 353.

PmpC(3) was amplified by the sense and antisense primers:

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GAGAGCGGCCGCTCGATCAATCTAACGAAAACACAGACG (SEQ ID NO: 354)
GAGAGCGGCCGCTTAGACCAAAGCTCCATCAGCAAC (SEQ ID NO: 355)
respectively. The resulting fusion construct has a DNA sequence set forth in SEQ ID

NO: 356 and encodes a 70 kD protein (683aa) expressing the segment 714-1250 aa of PmpC. The amino acid sequence of the fusion protein is set forth in SEO ID NO: 357.

EXAMPE 11

IMMUNOGENICITY OF CT622

Chlamydia-specific T cells lines were generated from two patients with

20 Chlamydia infections and the lines were designated CT1 and CT13. The T cell lines were either generated against monocyte-derived dendritic cells infected C. trachomatis serovar E for 72 hours (CT1-ERB) or against killed serovar E elementary bodies (EB) (CT13-EEB). Once generated, the lines were tested against the recombinant Chlamydia-specific protein, CT622 in a proliferation assay. Proliferation assays were performed by stimulating 2.5x10⁴ T cells in the presence of 1x10⁴ monocyte-derived dendritic cells with either recombinant CT antigens (2µg/ml) or Chlamydia EBs (1µg/ml). The assay was incubated for 4 days with a ³H-thymidine pulse for the last 18 hours.

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The cell line CT1-ERB demonstrated proliferative responses significantly above the media controls when stimulated with CT622, CT875, and CT EB. The cell line CT13-EEB demonstrated a proliferative response significantly above media controls when stimulated with CT622, CT875, and CT EB (see Figure 12).

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EXAMPLE 12

CLONING AND EXPRESSION OF FULL LENGTH CHLAMYDIA TRACHOMATIS GENES CT611, ORF3 AND OppA1

Recombinant protein expression of the full-length open reading frames was performed for clones containing genes CT611, ORF-3, and oppA1. The clones that contained the genes of interest were CtL2-8 (SEQ ID NO:285) which encoded 4 ORFs (CT474, CT473, CT060, and CT139), CtL2-10 (SEQ ID NO:284) which encoded the ORFs of CT610 and CT611, and clones 16CtL2-16 (SEQ ID NO:47), 16-D4-22 (SEQ ID NO:119) and 19-A5-54 (SEQ ID NO:84) which all contained sequences related to ORF-3. Sequences within CtL2-10 (Ct-610) and CtL2-16 (ORF-3) were also independently identified by the T-cell expression cloning approach. The clone CtL2-8 was further investigated as this clone had stimulated the proliferative responses and IFN-eamma production by two T cell lines generated against serovar E.

Cloning and expression of clone sequences:

Ctl.2-10 was found to encode two open reading frames (ORFs), CT610 and CT611, and these were found organized adjacent to each other within the genomic clone. The full length ORF of CT610 (containing a PQQ synthesis domain) was previously expressed and demonstrated to stimulate the proliferative responses of T cell lines generated against Chlamydia. To determine whether the second ORF, CT611, was also recognized by T cells, the full-length sequence of CT611 was PCR amplified and engineered for protein expression. The nucleotide sequence is disclosed in SEQ ID NO:365.

The second serological clone, Ctl.2-8, was found to contain 4 ORFs (CT474,
CT473, CT060, and CT139). Overlapping peptides to the three smallest predicted
ORFs (CT474, CT473, and CT060) did not stimulate the proliferative responses of T

cell lines. This suggested that the immunostimulatory antigen resides in the fourth ORF, CT139. The ORF of CT139 is approximately 450 nucleotides. The full-length nucleotide sequence is disclosed in SEQ ID NO:359 and the full-length amino acid sequence is disclosed in SEQ ID NO:363. Amino acid sequence comparison from 5 Genbank revealed that it is an oligo-peptide binding protein (oppA1) as well as belonging to the peptide ABC transporter family. This protein is 462 amino acids long with a predicted size of 48.3kDa and appears to contain 2 trans-membrane regions.

To express the full-length sequence of oppA1, oligonucleotides were designed which specifically amplified sequences starting from amino acid residue 22 (devoid of the first transmembrane domain), the nucleotide sequence for which is disclosed in SEQ ID NO:358 and, the amino acid sequence of which is disclosed in SEQ ID NO:362. This was shown to express the protein in E. coli.

The full-length cloning and recombinant protein expression of ORF-3 was also achieved. The nucleotide and amino acid sequences are disclosed in SEQ ID NOs:360

and 364, respectively.

EXAMPLE 13

RECOMBINANT CHLAMYDIAL ANTIGENS RECOGNIZED BY T CELL LINES

Patient T cell lines were generated from the following donors: CT1, CT2, CT3, CT4, CT5, CT6, CT7, CT8, CT9, CT10, CT11, CT12, CT13, CT14, CT15, and CT16, some of which were discussed above. A summary of their details is included in Table V.

	Table V: C. trachomatis patients											
Patients	Gender	Age	Clinical	Serovar	IgG	Multiple						
			Manifestation		titer	Infections						
CT1	M	27	NGU	LCR	Negative	No						
CT2	M	24	NGU	D	Negative	Е						
CT3	M	43	Asymptomatic	J	Ct 1:512	No						

		T	Shed Eb		Cp	
			Dx was HPV		1:1024	
		ĺ	ĺ		Cps	
					1:256	
CT4	F	25	Asymptomatic	J	Ct	Y
			Shed Eb		1:1024	E.
CT5	F	27	BV	LCR	Ct 1:256	F/F
				ļ	Ср	
					1:256	
CT6	M	26	Perinial rash	G	Ср	N
			Discharge,		1:1024	
			dysuria			
CT7	F	29	BV	Е	Ct 1:512	N
			Genital ulcer		Cp	
		l			1:1024	
CT8	F	24	Not Known	LCR	Not	NA
					tested	
CT9	M	24	asymptomatic	LCR	Ct 1:128	N
					Cp	
					1:128	
CT10	F	20	Mild itch vulvar	negative	negative	12/1/98
CT11	F	21	BV	J	Ct 1:512	F/F/J/E/E
			Abnormal pap			PID 6/96
			smear			
CT12	M	20	asymptomatic	LCR	Cp	N
					1:512	
CT13	F	18	BV, gonorrhea,	G	Ct	N
			Ct vaginal		1:1024	
			discharge, dysuria			
CT14	M	24	NGU	LCR	Ct 1:256	N

					Ср	
					1:256	
CT15	F	21	Muco-purulint	culture	Ct 1:256	N
			cervicitis		Ct IgM	
			Vaginal discharge		1:320	
					Cp 1:64	
CT16	M	26	Asymptomatic/	LCR	NA	N
			contact			
CL8	M	38	No clinical	negative	negative	N
			history of disease			

NGU=Non-Gonococcal Urethritis; BV=Bacterial Vaginosis; CT=Chlamydia trachomatis; Cp=Chlamydia pneumoniae; Eb=Chlamydia elementary bodies; HPV=human papiloma virus; Dx=diagnosis; PID=pelvic inflammatory disease; LCR=Ligase chain reaction.

PBMC were collected from a second series of donors and T cell lines have been generated from a sub-set of these. A summary of the details for three such T cell lines is listed in the table below.

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	Table III: Normal Donors											
Donor	Gender	Age	CT IgG Titer	CP IgG Titer								
CHH011	F	49	1:64	1:16								
СНН037	F	22	0	0								
CHH042	F	25	0	1:16								

Donor CHH011 is a heathly 49 year old female donor sero-negative for

C. trachomatis. PBMC produced higher quantities of IFN-gamma in response to C.

trachomatis elementary bodies as compared to C. pneumoniae elementary bodies,
indicating a C. trachomatis-specific response. Donor CHH037 is a 22 year old healthy

female donor sero-negative for C. trachomatis. PBMC poruced higher quantities of IFN-gamma in response to C. trachomatis elementary bodies as compared to C. pneumoniae elementary bodies, indicating a C. trachomatis-specific response. CHH042 is a 25 year old healthy female donor with an IgG titer of 1:16 to C. pneumoniae.

5 PBMC produced higher quantities of IFN-gamma in response to C. trachomatis elementary bodies as compared to C. pneumoniae elementary bodies, indicating a C. trachomatis-specific response.

Recombinant proteins for several *Chlamydia trachomatis* genes were generated as described above. Sequences for MOMP was derived from serovar F. The genes CT875, CT622, pmp-B-2, pmpA, and CT529 were derived from serovar E and sequences for the genes gro-EL, Swib, pmpD, pmpG, TSA, CT610, pmpC, pmpE, S13, lndA, pmpI, and pmpH-C were derived from LII.

Several of the patient and donor lines described above were tested against the recombinant Chlamydia proteins. Table IV summarizes the results of the T cell responses to these recombinant Chlamydia proteins.

	Table VII: Recombinant Chlamydia Antigens Recognized By T Cell Lines												
Antigen	Sero-	#of	С	CT	CT1	CT3	CT4	CT5	CT	CT	CT	CH	CH
	var	hits	L8	10	E	E	L2	E	11	12	13	H-	H-
			L2	E					E	Е	Е	011	037
												E	E
gro-EL (CT110)	L2	10	-	+	+	+	+	+	+	+	+	+	+
MompF (CT681)	F	10	-	+	+	+	+	+	+	+	+	+	+
CT875	Е	8	-	+	+	-	+	+	+	+	+	-	+
SWIB (CT460)	L2	8	+	+	-	+	-	+	-	+	+	+	+
pmpD (CT812)	L2	5	-	+	+	+	+	-	-	+	+	-	-

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pmpG (CT871)	L2	6	-	+	+	-	+	+	nt	-	+	+	-
TSA	L2										_		
(CT603)	1.2	6	-	-	+	+	+	+	-	-	+	-	+
CT622	E	3	-	-	+	-	+	-	-	-	+	-	-
CT610	1.2	3	-	+	-	+	-	-	-	+	-	-	-
pmpB-2 (CT413)	E	3	-	-	+	+	+	-	-	-	-	-	-
pmpC (CT414)	L2	4	-	-	-	+	-	+	-	+	-	-	+
pmpE (CT869)	L2	3	-	+	+	-	-	-	-	-	+	-	-
S13 (CT509)	L2	2	+	-	-	-	+	-	-	-	-	-	-
lpdA (CT557)	L2	3	-	-	+	+	-	-	-	-	-	+	-
pmpI (CT874)	L2	2	-	-	+	-	-	-	-	-	-	+	-
pmpH-C (CT872)	L2	1	-	•	-	-	-	-	-	+	-	-	-
pmpA (CT412)	Е	0	-	-	-	-	-	-	-	-	-	-	-
CT529	Е	0	-	-	-	-	-	-	-	-	-	-	-

Although the present invention has been described in some detail by way
of illustration and example for purposes of clarity of understanding, changes and
modifications can be carried out without departing from the scope of the invention
which is intended to be limited only by the scope of the appended claims.

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Claims

What is Claimed:

- A composition for eliciting an immune response comprising a Chlamydia Cap1
 protein or an immunogenic fragment thereof and an immunostimulant.
- The composition of claim 1, wherein the immunogenic fragment comprises at least a CTL epitope consisting essentially of amino acids 139-147 of a Cap1 protein.
- The composition of claim 1, wherein the Cap 1 protein comprises an amino acid sequence set forth in SEQ ID NO: 121 or a sequence having at least about 90% identity to the sequence set forth in SEO ID NO: 121.
- The composition of claim 1, wherein the Cap1 protein or immunogenic fragment thereof comprises a sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
- The composition of claim 1, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein.
- The composition of claim 5, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein having an amino acid sequence set forth in any one of SEO ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
 - 7. The composition of claim 1, wherein the immunogenic fragment is

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immunologically reactive with a CD8+ T-cell of a Chlamydia-infected animal.

- A method for stimulating a Chlamydia-specific T-cell response in an animal comprising administering to an animal an effective amount of a composition according to claim 1
- 9. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to an animal an effective amount of a composition according to claim 1.
- 10. A composition for eliciting an immune response comprising an isolated polynucleotide that encodes a Chlamydia Cap1 protein or an immunogenic fragment thereof and an immunostimulant.
- 11. The composition of claim 10, wherein the immunogenic fragment comprises at least the CTL epitope sequence consisting essentially of amino acids 139-147 of a Cap1 protein.
- 12. The composition of claim 10, wherein the Cap 1 protein comprises an amino acid sequence set forth in SEQ ID NO: 121 or a sequence having at least about 90% identity to the sequence set forth in SEQ ID NO: 121.
- 13. The composition of claim 10, wherein the Cap1 protein or immunogenic fragment thereof comprises a sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.

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- 14. The composition of claim 10, wherein the immunogenic fragment comprises amino acids 107-176 of a Cap1 protein.
- 15. The composition of claim 14, wherein the immunogenic fragment comprises amino acids 107-176 of a Capl protein having an amino acid sequence set forth in any one of SEQ ID NOs: 121, 123, 125, 127, 129, 131, 133, 135, 137 and 139.
- The composition of claim 10, wherein the immunogenic fragment is immunologically reactive with a CD8+ T-cell of a Chlamydia-infected animal.
- 17. The composition of claim 10, wherein the isolated polynucleotide is operably linked within a viral delivery vector.
- 18. The composition of claim 17, wherein the viral delivery vector is a vaccinia virus delivery vector.
- 19. A method for stimulating a Chlamydia-specific T-cell response in an animal comprising administering to said animal an effective amount of a composition according to claim 10.
- 20. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to an animal said effective amount of a composition according to claim 10.

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- 21. A method for inhibiting the development of a Chlamydia infection in an animal, comprising administering to said animal an effective amount of a composition according to claim 18.
- An isolated polynucleotide comprising a sequence selected from the group consisting of:
 - (a) sequences provided in SEQ ID NO:358-361;
 - (b) complements of the sequences provided in SEQ ID NO:358-361;
- (c) sequences consisting of at least 20 contiguous residues of a sequence provided in SEO ID NO:358-361;
- (d) sequences that hybridize to a sequence provided in SEQ ID NO:358-361, under highly stringent conditions;
- (e) sequences having at least 95% identity to a sequence of SEQ ID / NO:358-361;
- $\label{eq:sequences} \mbox{(f)} \qquad \mbox{sequences having at least 99\% identity to a sequence of SEQ ID} \\ \mbox{NO:358-361; and}$
- (g) degenerate variants of a sequence provided in SEQ ID NO:358-361.
- 23. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of:
 - (a) sequences encoded by a polynucleotide of claim 22;
- (b) sequences having at least 95% identity to a sequence encoded by a polynucleotide of claim 22; and
- (c) sequences having at least 99% identity to a sequence encoded by a polynucleotide of claim 22.
- 24. An isolated polypeptide comprising at least an immunogenic fragment of a polypeptide sequence selected from the group consisting of:
 - (a) a polypeptide sequence set forth in SEQ ID NO:362-365,
- (b) a polypeptide sequence having at least 95% identity with a sequence set forth in SEQ ID NO:362-365, and

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- (c) a polypeptide sequence having at least 99% identity with a sequence set forth in SEO ID NO:362-365.
- An expression vector comprising a polynucleotide of claim 22 operably linked to an expression control sequence.
- A host cell transformed or transfected with an expression vector according to claim 25.
- 27. An isolated antibody, or antigen-binding fragment thereof, that specifically binds to a polypeptide of any one of claims 23 and 24.
- 28. A method for detecting the presence of Chlamydia in a patient, comprising the steps of:
 - (a) obtaining a biological sample from the patient;
- (b) contacting the biological sample with a binding agent that binds to a polypeptide of any one of claims 23 and 24;
- (c) detecting in the sample an amount of polypeptide that binds to the binding agent; and
- (d) comparing the amount of polypeptide to a predetermined cut-off value and therefrom determining the presence of Chlamydia in the patient.
- A fusion protein comprising at least one polypeptide according to claim
 or claim 24.
- An oligonucleotide that hybridizes to a sequence recited in SEQ ID NO:
 358-361 under highly stringent conditions.
- 31. A method for stimulating and/or expanding T cells specific for a Chlamydia protein, comprising contacting T cells with at least one component selected from the group consisting of:
 - (a) a polypeptide according to claim 23 or claim 24;
 - (b) a polynucleotide according to claim 22; and

- (c) an antigen-presenting cell that expresses a polynucleotide according to claim 22,
- under conditions and for a time sufficient to permit the stimulation and/or expansion of T cells.
- 32. An isolated T cell population, comprising T cells prepared according to the method of claim 31.
- 33. A composition comprising a first component selected from the group consisting of physiologically acceptable carriers and immunostimulants, and a second component selected from the group consisting of:
 - (a) a polypeptide according to claim 23 or claim 24;
 - (b) a polynucleotide according to claim 22;
 - (c) an antibody according to claim 27;
 - (d) a fusion protein according to claim 29;
 - (e) a T cell population according to claim 32; and
- (f) an antigen presenting cell that expresses a polypeptide according to claim 23 or claim 24.
- 34. A method for stimulating an immune response in a patient, comprising administering to the patient a composition selected from the group consisting of;
 - (a) a composition of claim 33;
 - (b) a polynucleotide sequence of any one of SEQ ID NO:407-430, 525-559, and 582-598; and
 - (e) a polypeptide sequence of any one of SEQ ID NO:431-454 and 560-581.
- A method for the treatment of Chlamydia infection in a patient, comprising administering to the patient a composition selected from the group consisting of;
 - (a) a composition of claim 33;

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- (b) a polynucleotide sequence of any one of SEQ ID NO: 407-430,525-559, and 582-598; and
- (d) a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581.
- 36. A method for determining the presence of Chlamydia in a patient, comprising the steps of:
 - (a) obtaining a biological sample from the patient;
- (c) detecting in the sample an amount of a polynucleotide that hybridizes to the oligonucleotide; and
- (d) comparing the amount of polynucleotide that hybridizes to the oligonucleotide to a predetermined cut-off value, and therefore determining the presence of the cancer in the patient.
- A diagnostic kit comprising at least one oligonucleotide according to claim 30.
- 38. A diagnostic kit comprising at least one antibody according to claim 27 and a detection reagent, wherein the detection reagent comprises a reporter group.
- 39. A method for the treatment of Chlamydia in a patient, comprising the steps of:
- (a) incubating CD4+ and/or CD8+ T cells isolated from a patient with at least one component selected from the group consisting of:
 - (i) a polypeptide according to any one of claims 23 and 24;
 - (ii) a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581;
 - (iii) a polynucleotide according to claim 22;
 - (iv) a polynucleotide sequence of any one of SEQ ID NO: 407-430, 525-559 and 582-598;

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 an antigen presenting cell that expresses a polypeptide sequence set forth in any one of claims 23 and 24;

- (vi) an antigen presenting cell that expresses a polypeptide sequence of any one of SEQ ID NO: 431-454 and 560-581, such that the T cells proliferate; and
- (b) administering to the patient an effective amount of the proliferated T cells.

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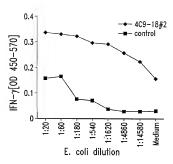


Fig. 1

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Fig. 2

Chlamydia C17.8 Peptide Screen

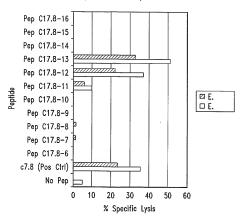
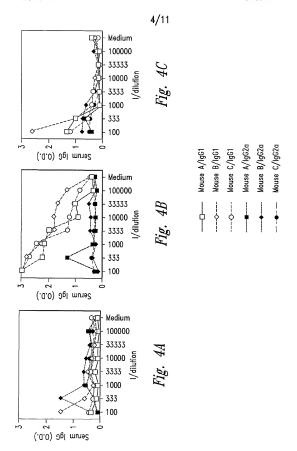
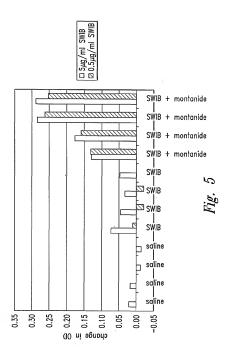


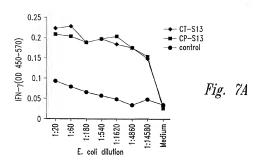
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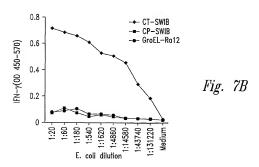




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- CP SWIB EcoRI (3' primer)
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Fig. 6





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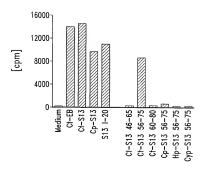
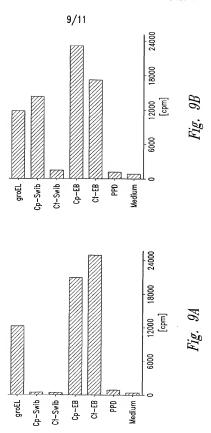


Fig. 8





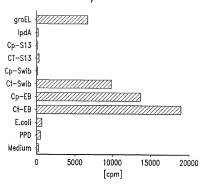


Fig. 10

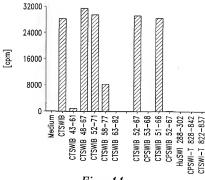


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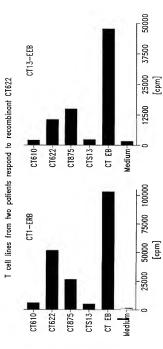


Fig. 15

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<400> 26

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<400> 27

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tectatette ticagetata assatzette tiassactie atateeteta atcasatcat 180
cattaaccac aacataatca aattegetag eggeageaat ttegacageg etatgeteta 240
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Phe Lys Ile Ser Glu Ala Phe Gly Val Leu Asn Pro Glu Gly Ser Leu 115 \$120\$

Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg His

Ala Val Ile Asn Asp Leu Pro Leu Gly Arg Ser Ile Asp Glu Glu Leu

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WO 02/08267 PCT/US01/23121 36

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300

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420

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660

720

780

840

897

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<210> 127 <211> 298

<212> PRT <213> Chlamydia

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540

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660

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<400> 128

<210> 129 <211> 298

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280

<210> 130 <211> 897

275

<212> DNA

<213> Chlamydia

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<400> 130

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285

180

300

360

420

480

660

720

780

840

897

60

120

agctatatta tggcggctaa ccatgcagcg tttgtggtgg gttctggact cgctatcagt geggaaagag cagattgega agecegetge getegtattg egagagaaga gtegteacte gaattgtegg gagaggaaaa tgettgegag aggggagteg etggagagaa agecaagaeg ttcacgcgca tcaagtatgc actcctcact atgctcgaga agtttttgga atgcgttgcc gacgittica aattggtgcc gttgcctatt acaatgggta ttcgtgcaat tgtggctgcg ggatgtacgt tcacttctgc agttattgga ttgtggactt tctgcaacag agtataa <210> 131 <211> 298 <212> PRT <213> Chlamydia <400> 131 Met Ala Ala Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu Lys Ala Phe Phe Thr Gln Pro Ser Asn Lys Met Ala Arg Val Val Asn 25 Lys Thr Lys Gly Met Asp Lys Thr Val Lys Val Ala Lys Ser Ala Ala Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Ala Arg 7.0 Thr Val Leu Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr Val Gln Ser Ala Gln Ser Phe Phe Ser Tyr Met Lys Ala Ala Ser Gln 100 105 Lys Pro Gln Glu Gly Asp Glu Gly Leu Val Ala Asp Leu Cys Val Ser 120 His Lys Arg Arg Ala Ala Ala Ala Val Cys Ser Phe Ile Gly Gly Ile 135 140 Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn 150 155 Lys Met Leu Ala Gln Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met 165 170 Gly Ser Ser Val Ser Tyr Ile Met Ala Ala Asn His Ala Ala Phe Val 180 185 Val Gly Ser Gly Leu Ala Ile Ser Ala Glu Arg Ala Asp Cys Glu Ala 195 205 200 Arg Cys Ala Arg Ile Ala Arg Glu Glu Ser Ser Leu Glu Leu Ser Gly 210 215 Glu Glu Asn Ala Cys Glu Arg Gly Val Ala Gly Glu Lys Ala Lys Thr 230 235 Phe Thr Arg Ile Lys Tyr Ala Leu Leu Thr Met Leu Glu Lys Phe Leu 250 Glu Cys Val Ala Asp Val Phe Lys Leu Val Pro Leu Pro Ile Thr Met 260 265 270 Gly Ile Arg Ala Ile Val Ala Ala Gly Cys Thr Phe Thr Ser Ala Val 275 280 Ile Gly Leu Trp Thr Phe Cys Asn Arg Val 290 295

<210> 132 <211> 897

<212> DNA <213> Chlamydia

<400> 132

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300

360

420

480

540

600

660

720 780

840

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caaagettet tetettacat gaaagetget agteagaaac egcaagaagg ggatgagggg
ctcgtagcag atctttgtgt gtctcataag cgcagagcgg ctgcggctgt ctgtagcttc
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aaaatgetgg cgcaaccgtt tetttettee caaactaaag caaatatggg atettetgtt
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gaattgtcqq qaqaqqaaaa tqcttgtgag aggaqaqtcq ctqqagaqaa agccaagacq
ttcacgcgca tcaagtatgc actcctcact atgctcgaga agtttttgga atgcgttgcc
gacqttttca aattggtgcc gttgcctatt acaatgggta ttcgtgcaat tgtggctgcg
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Lys Thr Lys Gly Met Asp Lys Thr Val Lys Val Ala Lys Ser Ala Ala
                          40
Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser
Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Ala Arg
                   70
Thr Val Leu Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr
Val Gln Ser Ala Gln Ser Phe Phe Ser Tyr Met Lys Ala Ala Ser Gln
                               105
Lys Pro Gln Glu Gly Asp Glu Gly Leu Val Ala Asp Leu Cys Val Ser
                           120
His Lys Arg Arg Ala Ala Ala Ala Val Cys Ser Phe Ile Gly Gly Ile
                       135
                                           140
Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
                   150
                                       155
Lys Met Leu Ala Gln Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met
               165
                                   170
Gly Ser Ser Val Ser Tyr Ile Met Ala Ala Asn His Ala Ala Phe Val
           180
                               185
Val Gly Ser Gly Leu Ala Ile Ser Ala Glu Arg Ala Asp Cys Glu Ala
                           200
Arg Cys Ala Arg Ile Ala Arg Glu Glu Ser Ser Leu Glu Leu Ser Gly
                       215
Glu Glu Asn Ala Cys Glu Arg Arg Val Ala Gly Glu Lys Ala Lys Thr
                                       235
                   230
Phe Thr Arg Ile Lys Tyr Ala Leu Leu Thr Met Leu Glu Lys Phe Leu
               245
                                  250
Glu Cys Val Ala Asp Val Phe Lys Leu Val Pro Leu Pro Ile Thr Met
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Gly Ile Arg Ala Ile Val Ala Ala Gly Cys Thr Phe Thr Ser Ala Val
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                           280
Ile Gly Leu Trp Thr Phe Cys Asn Arg Val
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420

480

540

600

660

780

840

897

<213> Chlamydia

<400> 134

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<210> 135

<211> 298 <212> PRT

<213> Chlamydia

<400> 135

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780

840

882

Ile Gly Leu Cys Thr Phe Cys Ala Arg Ala 290 <210> 136 <211> 882 <212> DNA <213> Chlamydia <400> 136 atggettetg tatgtgggeg attaagtget ggggtgggga acagatttaa egcattttte acgcgtcccg gtaacaaget atcacggttt gtaaatagcg caaaaggatt agacagatca 120 ataaaggttg ggaagtctgc tgctgaatta acggcgagta ttttagagca aactgggggg 180 geagggactg atgeacatgt taeggeggee aaggtgteta aageacttgg ggaegegega 240 acagtaatgg ctctagggaa tgtcttcaat gggtctgtgc cagcaaccat tcaaagtgcg egaagetgte tegeceattt acqageggee ggcaaagaag aagaaacatg etecaaggtg aaagatetet gtgtttetea tagaegaaga getgeggetg aggettgtaa tgttattgga 420 ggagcaactt atattacaac tttcggagcg attcgtccga cattactcgt taacaagctt 480 ettgecaaac catteettte eteccaagee aaagaagggt tgggagette tgttggttat 540 atcatggcag cgaaccatgc ggcatctgtg cttgggtctg ctttaagtat tagegcagaa 600 agagcagact gtgaagagcg gtgtgatcgc attcgatgta gtgaggatgg tgaaatttgc

gaaggcaata aattaacagc tattteggaa gagaaggeta gatcatggac teteattaag

tacagattee ttactatgat agaaaaacta tttgagatgg tggeggatat etteaagtta

attecttige caattiegea tggaattegt getattgitg etgegggatg taegttgaet

totgoagtta ttggottagg tactttttgg totagagcat aa

<210> 137

<211> 293 <212> PRT

<213> Chlamydia

<400> 137

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Tyr Arg Phe Leu Thr Met Ile Glu Lys Leu Phe Glu Met Val Ala Asp
              245
                           250
Ile Phe Lys Leu Ile Pro Leu Pro Ile Ser His Gly Ile Arg Ala Ile
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Val Ala Ala Gly Cys Thr Leu Thr Ser Ala Val Ile Gly Leu Gly Thr
275 280 285
Phe Trp Ser Arg Ala
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Arg Ala Ala Ala Val Cys Ser Phe Ile Gly Gly Ile Thr Tyr Leu
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      <211> 18
      <212> PRT
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     <220>
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Cys Ser Phe Ile Gly Gly Ile Thr Tyr Leu Ala Thr Phe Gly Ala Ile
Arg Pro
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Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn Lys
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Met Leu
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Ser Gln
     <210> 143
     <211> 17
     <212> PRT
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     <400> 143
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Ser
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Cys Ser Phe Ile Gly Gly Ile Thr Tyr Leu
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Ser Phe Ile Gly Gly Ile Thr Tyr Leu
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     <400> 147
Cys Ser Phe Ile Gly Gly Ile Thr Tyr
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     <211> 8
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     <220>
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Cys Ser Phe Ile Gly Gly Ile Thr
     <210> 149
      <211> 10
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     <223> Made in a lab
     <400> 149
Cys Ser Ile Ile Gly Gly Ile Thr Tyr Leu
     <210> 150
<211> 10
      <212> PRT
     <213> Artificial Sequence
     <220>
     <223> Made in a lab
     <400> 150
Cys Gly Phe Ile Gly Gly Ile Thr Tyr Leu
     <210> 151
     <211> 9
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     <223> Made in a lab
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Gly Phe Ile Gly Gly Ile Thr Tyr Leu
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     <220>
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     <400> 152
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Ser Val Ala Ser
           20
     <210> 153
     <211> 20
     <212> PRT
     <213> Artificial Sequence
     <220>
     <223> Made in a lab
     <400> 153
Glu Arg Leu Arg Leu Arg Leu Ser Val Ala Ser Ser Glu Glu Leu Pro
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Thr Ser Arg His
           20
     <210> 154
     <211> 20
     <212> PRT
     <213> Artificial Sequence
     <220>
     <223> Made in a lab
    <400> 154
Ala Ser Ser Glu Glu Leu Pro Thr Ser Arg His Ser Glu Leu Ser Val
1
                                  10
Arg Phe Cys Leu
           20
     <210> 155
     <211> 20
     <212> PRT
     <213> Artificial Sequence
     <220>
     <223> Made in a lab
     <400> 155
Arg His Ser Glu Leu Ser Val Arg Phe Cys Leu Ser Thr Lys Cys Trp
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1
Arg Asn Arg Phe
           20
     <210> 156
     <211> 20
     <212> PRT
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52

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Gln Ile Trp Asp
     <210> 157
     <211> 53
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     <220>
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     <400> 157
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1.
                                 10
                                                  15
Ser Ser Glu Glu Leu Pro Thr Ser Arg His Ser Glu Leu Ser Val Arg
        20
                                                30
                             25
Phe Cys Leu Ser Thr Lys Cys Trp Arg Asn Arg Phe Phe Leu Pro Lys
       35
                          40
Leu Lys Gln Ile Trp
   50
     <210> 158
     <211> 52
     <212> PRT
     <213> Artificial Sequence
     <220>
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                                 10
                                                   15
Ile Gly Gly Ile Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile
           20
                              25
                                                30
Leu Phe Val Asn Lys Met Leu Ala Gln Pro Phe Leu Ser Ser Gln Ile
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                                             45
Lvs Ala Asn Met
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     <211> 24
     <212> DNA
     <213> Chlamydia
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     <210> 160
     <211> 24
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<400> 160

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<210> 163 <211> 24 <212> DNA <213> Chlamydia		
<400> 163 ttttgaagca ggtaggtgaa	tatg	24
<210> 164 <211> 29 <212> DNA <213> Chlamydia		
<400> 164 tttacaataa gaaaagctaa	gcactttgt	29
<210> 165 <211> 20 <212> DNA <213> Chlamydia		
<400> 165 ccttacacag tcctgctgac		20
<210> 166 <211> 20 <212> DNA <213> Chlamydia		
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<210> 167 <211> 9 <212> PRT <213> Artificia	1 Sequence	
<220> <223> Made in a	lab	
<400> 167 Ser Phe Ile Gly Gly II 1 5	le Thr Tyr Leu	

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       <211> 9
       <212> PRT
       <213> Artificial Sequence
       <220>
       <223> Made in a lab
       <400> 168
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<210> 169
<211> 2643
<212> DNA
<213> Chlamydia
<400> 169
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acageggtee tetttggeca ggateeetta ggtgaaaceg ceeteeteac taaaaateet
                                                                      120
aatcatgtcg totgtacatt ttttgaggac tgtaccatgg agagcotott tootgotott
                                                                      180
tgtgctcatg catcacaaga cgatcctttg tatgtacttg gaaattccta ctgttggttc
                                                                      240
gtatetaaae tecatateae ggaceecaaa gaggetettt ttaaagaaaa aggagatett
                                                                      300
tocattoaaa actitogott cotttoctto acagattgot ottocaagga aagetotoot
                                                                      360
totattatto atcassagas tggtcagtta toottgcgca ataatggtag catgagttto
                                                                      420
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<213> Chlamydia

<220> <221> VARIANT

<222> 336

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63

645 650 Ala Ala Gly Gln Leu Leu Gly Lys Ser Ser Asp Ser Phe Ile Thr Ser 665 660 67.0 Thr Glu Thr Thr Ser Tyr Ile Ala Thr Val Gln Ala Gln Leu Ala Thr 680 685 Ser Leu Met Lys Ile Ser Ala Gln Ala Cys Tyr Asn Glu Ser Ile His 695 700 Glu Leu Lys Thr Lys Tyr Arg Ser Phe Ser Lys Glu Gly Phe Gly Ser 705 $\,$ 710 $\,$ 720 Trp His Ser Val Ala Val Ser Gly Glu Val Cys Ala Ser Ile Pro Ile 725 730 735 Val Ser Asn Gly Ser Gly Leu Phe Ser Ser Phe Ser Ile Phe Ser Lys Leu Gln Gly Phe Ser Gly Thr Gln Asp Gly Phe Glu Glu Ser Ser Gly 760 765 Glu Ile Arg Ser Phe Ser Ala Ser Ser Phe Arg Asn Ile Ser Leu Pro 775 780 Ile Gly Ile Thr Phe Glu Lys Lys Ser Gln Lys Thr Arg Thr Tyr Tyr 790 795 Tyr Phe Leu Gly Ala Tyr Ile Gln Asp Leu Lys Arg Asp Val Glu Ser 805 810 Gly Pro Val Val Leu Leu Lys Asn Ala Val Ser Trp Asp Ala Pro Met 820 825 830Ala Asn Leu Asp Ser Arg Ala Tyr Met Phe Arg Leu Thr Asn Gln Arg 835 840 845 Ala Leu His Arg Leu Gln Thr Leu Leu Asn Val Ser Cys Val Leu Arg 855 860 Gly Gln Ser His Ser Tyr Ser Leu Asp Leu Gly Thr Thr Tyr Arg Phe 875 <210> 176 <211> 982 <212> PRT <213> Chlamydia <220> <221> VARIANT <222> 981 <223> Xaa = Any Amino Acid Met Ile Pro Gln Gly Ile Tyr Asp Gly Glu Thr Leu Thr Val Ser Phe 5 10 Pro Tyr Thr Val Ile Gly Asp Pro Ser Gly Thr Thr Val Phe Ser Ala 25 Gly Glu Leu Thr Leu Lys Asn Leu Asp Asn Ser Ile Ala Ala Leu Pro 35 40 45 Leu Ser Cys Phe Gly Asn Leu Leu Gly Ser Phe Thr Val Leu Gly Arg 55 Gly His Ser Leu Thr Phe Glu Asn Ile Arg Thr Ser Thr Asn Gly Ala 70 75 Ala Leu Ser Asn Ser Ala Ala Asp Gly Leu Phe Thr Ile Glu Gly Phe 85 90 Lys Glu Leu Ser Phe Ser Asn Cys Asn Ser Leu Leu Ala Val Leu Pro Ala Ala Thr Thr Asn Lys Gly Ser Gln Thr Pro Thr Thr Thr Ser Thr 115 120 125 Pro Ser Asn Gly Thr Ile Tyr Ser Lys Thr Asp Leu Leu Leu Leu Asn 130 135 140 Asn Glu Lys Phe Ser Phe Tyr Ser Asn Leu Val Ser Gly Asp Gly Gly

150

Ala Ile Asp Ala Lys Ser Leu Thr Val Gln Gly Ile Ser Lys Leu Cys 165 170 Val Phe Gln Glu Asn Thr Ala Gln Ala Asp Gly Gly Ala Cys Gln Val 180 185 Val Thr Ser Phe Ser Ala Met Ala Asn Glu Ala Pro Ile Ala Phe Val 195 200 205 Ala Asn Val Ala Gly Val Arg Gly Gly Gly Ile Ala Ala Val Gln Asp 210 215 220 Gly Gln Gln Gly Val Ser Ser Ser Thr Ser Thr Glu Asp Pro Val Val 225 230 235 240 Ser Phe Ser Arg Asn Thr Ala Val Glu Phe Asp Gly Asn Val Ala Arg 250 Val Gly Gly Gly Tle Tyr Ser Tyr Gly Asn Val Ala Phe Leu Asn Asn 260 265 270 265 Gly Lys Thr Leu Phe Leu Asn Asn Val Ala Ser Pro Val Tyr Ile Ala 275 280 285 Ala Lys Gln Pro Thr Ser Gly Gln Ala Ser Asn Thr Ser Asn Asn Tyr 290 295 300 Gly Asp Gly Gly Ala Ile Phe Cys Lys Asn Gly Ala Gln Ala Gly Ser 305 310 315Asn Asn Ser Gly Ser Val Ser Phe Asp Gly Glu Gly Val Val Phe Phe 325 330 335 Ser Ser Asn Val Ala Ala Gly Lys Gly Gly Ala Ile Tyr Ala Lys Lys 340 345 350 Leu Ser Val Ala Asn Cys Gly Pro Val Gln Phe Leu Arg Asn Tle Ala 355 360 365 Asn Asp Gly Gly Ala Ile Tyr Leu Gly Glu Ser Gly Glu Leu Ser Leu 370 375 380 Ser Ala Asp Tyr Gly Asp Ile Ile Phe Asp Gly Asn Leu Lys Arg Thr 385 390 395 400 Ala Lys Glu Asn Ala Ala Asp Val Asn Gly Val Thr Val Ser Ser Gln 405 410 Ala Ile Ser Met Gly Ser Gly Gly Lys Ile Thr Thr Leu Arg Ala Lys Ala Gly His Gln Tle Leu Phe Asn Asp Pro Tle Glu Met Ala Asn Gly 435 440 445 Asn Asn Gln Pro Ala Gln Ser Ser Lys Leu Leu Lys Ile Asn Asp Gly $450 \ \ \, 455 \ \ \, 460 \ \ \,$ Glu Gly Tyr Thr Gly Asp Ile Val Phe Ala Asn Gly Ser Ser Thr Leu 465 470 475 480 Tyr Gln Asn Val Thr Ile Glu Gln Gly Arg Ile Val Leu Arg Glu Lys 485 490 495Ala Lys Leu Ser Val Asn Ser Leu Ser Gln Thr Gly Gly Ser Leu Tyr
500 505 510 Met Glu Ala Gly Ser Thr Leu Asp Phe Val Thr Pro Gln Pro Pro Gln 515 520 525 Gln Pro Pro Ala Ala Asn Gln Leu Ile Thr Leu Ser Asn Leu His Leu 535 540 Ser Leu Ser Ser Leu Leu Ala Asn Asn Ala Val Thr Asn Pro Pro Thr 545 550 555 560550 Asn Pro Pro Ala Gln Asp Ser His Pro Ala Val Ile Gly Ser Thr Thr 565 570 575 Ala Gly Ser Val Thr Ile Ser Gly Pro Ile Phe Phe Glu Asp Leu Asp 580 585 590Asp Thr Ala Tyr Asp Arg Tyr Asp Trp Leu Gly Ser Asn Gln Lys Ile 595 600 605 Asn Val Leu Lys Leu Gln Leu Gly Thr Lys Pro Pro Ala Asn Ala Pro 610 615 620 Ser Asp Leu Thr Leu Gly Asn Glu Met Pro Lys Tyr Gly Tyr Gln Gly 625 630 635 Ser Trp Lys Leu Ala Trp Asp Pro Asn Thr Ala Asn Asn Gly Pro Tyr

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<400> 177

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Phe	Glu 130	Asn	Asn	Thr	Сув	Cys 135		Leu	Phe	Thr	Trp		Asn	Pro	Tyr
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	210				Gly	215					220				
225					Cys 230					235					240
WIA	Asn	_		245	Phe				250				_	255	
GIA			260		Tyr			265			-		270		
		275			Asn	-	280	_			-	285			•
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Gly	Ala	Ile	Tyr	325 Ile	Asp				330					335	_
Arg	His		340 Ile	Ile	Phe	Asn		345 Asn	Ile	Val	Thr		350 Val	Thr	Asn
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Ser	Gly	Ala 435		Val	Asn	Ser	Ala		Phe	His	Gln	Arg		Leu	Gln
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Val				485	Asn				490					495	
Thr			500		Ser			505					510		-
		515			Ile		520					525			
	530				Asn	535				-	540				
545					Ser 550	-		-		555			-	-	560
eTA	Asn	ser	PTO	565	Glu	ser	rnr	Asp	Leu 570	Thr	HIS	Ата	ьeu	Ser 575	Ser

Gln Pro Met Leu Ser Ile Ser Glu Ala Ser Asp Asn Gln Leu Gln Ser 580 585 Glu Asn Ile Asp Phe Ser Gly Leu Asn Val Pro His Tyr Gly Trp Gln 600 Gly Leu Trp Thr Trp Gly Trp Ala Lys Thr Gln Asp Pro Glu Pro Ala 610 615 620 Ser Ser Ala Thr Ile Thr Asp Pro Gln Lys Ala Asn Arg Phe His Arg 625 630 635 640 Thr Leu Leu Leu Thr Trp Leu Pro Ala Gly Tyr Val Pro Ser Pro Lys 645 650 655 His Arg Ser Pro Leu Ile Ala Asn Thr Leu Trp Gly Asn Met Leu Leu. 660 665 670 Ala Thr Glu Ser Leu Lys Asn Ser Ala Glu Leu Thr Pro Ser Gly His 680 Pro Phe Trp Gly Ile Thr Gly Gly Gly Leu Gly Met Met Val Tyr Gln Asp Pro Arg Glu Asn His Pro Gly Phe His Met Arg Ser Ser Gly Tyr 705 710 715 720 Ser Ala Gly Met Ile Ala Gly Gln Thr His Thr Phe Ser Leu Lys Phe 725 730 735 Ser Gln Thr Tyr Thr Lys Leu Asn Glu Arg Tyr Ala Lys Asn Asn Val 740 745 750 Ser Ser Lys Asn Tyr Ser Cys Gln Gly Glu Met Leu Phe Ser Leu Gln 755 760 765 Glu Gly Phe Leu Leu Thr Lys Leu Val Gly Leu Tyr Ser Tyr Gly Asp 770 775 780 His Asn Cys His His Phe Tyr Thr Gln Gly Glu Asn Leu Thr Ser Gln 785 790 800 Gly Thr Phe Arg Ser Gln Thr Met Gly Gly Ala Val Phe Phe Asp Leu 805 810 815 Pro Met Lys Pro Phe Gly Ser Thr His Ile Leu Thr Ala Pro Phe Leu 820 825 830 Gly Ala Leu Gly Ile Tyr Ser Ser Leu Ser His Phe Thr Glu Val Gly 835 840 845 Ala Tyr Pro Arg Ser Phe Ser Thr Lys Thr Pro Leu Ile Asn Val Leu 850 855 860 Val Pro Ile Gly Val Lys Gly Ser Phe Met Asn Ala Thr His Arg Pro 865 870 875 Gln Ala Trp Thr Val Glu Leu Ala Tyr Gln Pro Val Leu Tyr Arg Gln 885 890 895 Glu Pro Gly Ile Ala Thr Gln Leu Leu Ala Ser Lys Gly Ile Trp Phe $900 \hspace{1.5cm} 905 \hspace{1.5cm} 910$ Gly Ser Gly Ser Pro Ser Ser Arg His Ala Met Ser Tyr Lys Ile Ser 915 920 925 Gln Gln Thr Gln Pro Leu Ser Trp Leu Thr Leu His Phe Gln Tyr His 935 940 Gly Phe Tyr Ser Ser Ser Thr Phe Cys Asn Tyr Leu Asn Gly Glu Ile 950 955 Ala Leu Arg Phe

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<212> PRT <213> Chlamydia

<213> Chlamydia

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515 520 Gly Ile Ser Phe Thr Gly Asn Ala Arg Ala Pro Gln Ala Leu Pro Thr 530 535 540 Gln Glu Glu Phe Pro Leu Phe Ser Lys Lys Glu Gly Arg Pro Leu Ser 550 555 Ser Gly Tyr Ser Gly Gly Gly Ala Ile Leu Gly Arg Glu Val Ala Ile 565 570 575 Leu His Asn Ala Ala Val Val Phe Glu Gln Asn Arg Leu Gln Cys Ser 585 590 580 Glu Glu Glu Ala Thr Leu Leu Gly Cys Cys Gly Gly Gly Ala Val His 595 600 605 Gly Met Asp Ser Thr Ser Ile Val Gly Asn Ser Ser Val Arg Phe Gly 610 615 620 Asn Asn Tyr Ala Met Gly Gln Gly Val Ser Gly Gly Ala Leu Leu Ser 625 630 635 640 Lys Thr Val Gln Leu Ala Gly Asn Gly Ser Val Asp Phe Ser Arg Asn 645 650 655 Ile Ala Ser Leu Gly Gly Gly Ala Leu Gln Ala Ser Glu Gly Asn Cys 660 665 670 Glu Leu Val Asp Asn Gly Tyr Val Leu Phe Arg Asp Asn Arg Gly Arg 675 680 685 Val Tyr Gly Gly Ala Ile Ser Cys Leu Arg Gly Asp Val Val Ile Ser 690 695 700 Gly Asn Lys Gly Arg Val Glu Phe Lys Asp Asn Ile Ala Thr Arg Leu 705 710 715 720 Tyr Val Glu Glu Thr Val Glu Lys Val Glu Glu Val Glu Pro Ala Pro
725 730 735 Glu Gln Lys Asp Asn Asn Glu Leu Ser Phe Leu Gly Ser Val Glu Gln
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Thr Leu Cys Ser Tyr Gly Phe Lys Gln Asp Ala Gly Ala Lys Leu Val Leu Ala Ala Gly Ser Lys Leu Lys Ile Leu Asp Ser Gly Thr Pro Val 1025 1030 1035 1040 Gln Gly His Ala Ile Ser Lys Pro Glu Ala Glu Ile Glu Ser Ser Ser 1045 1050 1055 Glu Pro Glu Gly Ala His Ser Leu Trp Ile Ala Lys Asn Ala Gln Thr 1060 1065 1070Thr Val Pro Met Val Asp Ile His Thr Ile Ser Val Asp Leu Ala Ser 1075 1080 1085 Phe Ser Ser Ser Gln Gln Glu Gly Thr Val Glu Ala Pro Gln Val Ile 1090 1095 1100 Val Pro Gly Gly Ser Tyr Val Arg Ser Gly Glu Leu Asn Leu Glu Leu 1105 1110 1115 1120 Val Asn Thr Thr Gly Thr Gly Tyr Glu Asn His Ala Leu Leu Lys Asn 1125 1130 135 Glu Ala Lys Val Pro Leu Met Ser Phe Val Ala Ser Ser Asp Glu Ala 1140 1145 1150Ser Ala Glu Ile Ser Asn Leu Ser Val Ser Asp Leu Gln Ile His Val 1155 1160 1165 Ala Thr Pro Glu Ile Glu Glu Asp Thr Tyr Gly His Met Gly Asp Trp 1170 1175 1180 Ser Glu Ala Lys Ile Gln Asp Gly Thr Leu Val Ile Asn Trp Asn Pro 1185 1190 1195 1200 Thr Gly Tyr Arg Leu Asp Pro Gln Lys Ala Gly Ala Leu Val Phe Asn 1205 1210 1215 Ala Leu Trp Glu Glu Gly Ala Val Leu Ser Ala Leu Lys Asn Ala Arg 1220 1225 1230 Phe Ala His Asn Leu Thr Ala Gln Arg Met Glu Phe Asp Tyr Ser Thr 1235 1240 1245 Asn Val Trp Gly Phe Ala Phe Gly Gly Phe Arg Thr Leu Ser Ala Glu 1250 1255 1260 Asn Leu Val Ala Ile Asp Gly Tyr Lys Gly Ala Tyr Gly Gly Ala Ser 1265 1270 1275 128 Ala Gly Val Asp Ile Gln Leu Met Glu Asp Phe Val Leu Gly Val Ser 1285 1290 1295 Gly Ala Ala Phe Leu Gly Lys Met Asp Ser Gln Lys Phe Asp Ala Glu 1300 1305 1310 Val Ser Arg Lys Gly Val Val Gly Ser Val Tyr Thr Gly Phe Leu Ala 1315 1320 1325 Gly Ser Trp Phe Phe Lys Gly Gln Tyr Ser Leu Gly Glu Thr Gln Asn 1330 1335 1340 Asp Met Lys Thr Arg Tyr Gly Val Leu Gly Glu Ser Ser Ala Ser Trp 1345 1350 1355 1360 Thr Ser Arg Gly Val Leu Ala Asp Ala Leu Val Glu Tyr Arg Ser Leu 1365 1370 1375 Val Gly Pro Val Arg Pro Thr Phe Tyr Ala Leu His Phe Asn Pro Tyr 1380 1385 1390 Val Glu Val Ser Tyr Ala Ser Met Lys Phe Pro Gly Phe Thr Glu Gln 1395 1400 1405 Gly Arg Glu Ala Arg Ser Phe Glu Asp Ala Ser Leu Thr Asn Ile Thr 1410 1415 1420 Ile Pro Leu Gly Met Lys Phe Glu Leu Ala Phe Ile Lys Gly Gln Phe 1425 1430 1435 1440 Ser Glu Val Asn Ser Leu Gly Ile Ser Tyr Ala Trp Glu Ala Tyr Arg 1445 1450 1450 Lys Val Glu Gly Gly Ala Val Gln Leu Leu Glu Ala Gly Phe Asp Trp 1460 1465 1470 Glu Gly Ala Pro Met Asp Leu Pro Arg Gln Glu Leu Arg Val Ala Leu 1475 1480 1485 Glu Asn Asn Thr Glu Trp Ser Ser Tyr Phe Ser Thr Val Leu Gly Leu

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Gly Ala Tyr Val Thr Gln Thr Met Ser Val Thr Asn Thr Thr Ser Glu

72

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Thr	Glu	Thr 515		Asp	Thr	Asn	Ser 520		Ile	Asp	Val	Ser 525		Glu	Asn
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Thr	Gly	Thr 675	Gly	Val	Val	Asn	Asn 680	Glu	Ser	Gln	Asp	Thr 685	Ser	Asp	Thr
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Ala	Cys 770	Leu	Ala	Lys	Ser	Tyr 775	Ala	Ala	Ser	Thr	Asp 780	Ser	Ser	Pro	Val
Ser 785	Asn	Ser	Ser	Gly	Ser 790	Asp	Val	Thr	Ala	Ser 795	Ser	Asp	Asn	Pro	Asp 800
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Pro	Glu	Ala	Gly 820	Ser	Thr	Thr	Glu	Thr 825	Pro	Thr	Leu	Ile	Gly 830	Gly	Gly
Ala	Ile	Tyr 835	Gly	Glu	Thr	Val	Lys 840		Glu	Asn	Phe	Ser 845	Gly	Gln	Gly
Ile	Phe 850	Ser	Gly	Asn	Lys	Ala 855		Asp	Asn	Thr	Thr 860	Glu	Gly	Ser	Ser
Ser 865	Lys	Ser	Asn	Val	Leu 870		Gly	Ala	Val	Tyr 875		Lys	Thr	Leu	Phe 880

Asn Leu Asp Ser Gly Ser Ser Arg Arg Thr Val Thr Phe Ser Gly Asn 890 Thr Val Ser Ser Gln Ser Thr Thr Gly Gln Val Ala Gly Gly Ala Ile 905 Tyr Ser Pro Thr Val Thr Ile Ala Thr Pro Val Val Phe Ser Lys Asn 915 920 925 Ser Ala Thr Asn Asn Ala Asn Asn Ala Thr Asp Thr Gln Arg Lys Asp 930 935 940 Thr Phe Gly Gly Ala Ile Gly Ala Thr Ser Ala Val Ser Leu Ser Gly 945 950 955 960 Gly Ala His Phe Leu Glu Asn Val Ala Asp Leu Gly Ser Ala Ile Gly 965 970 975 Leu Val Pro Asp Thr Gln Asn Thr Glu Thr Val Lys Leu Glu Ser Gly 980 985 Ser Tyr Tyr Phe Glu Lys Asn Lys Ala Leu Lys Arg Ala Thr Ile Tyr 995 1000 1005Ala Pro Val Val Ser Ile Lys Ala Tyr Thr Ala Thr Phe Asn Gln Asn 1010 1015 1020 1015 Arg Ser Leu Glu Glu Gly Ser Ala Ile Tyr Phe Thr Lys Glu Ala Ser 1025 1030 1035 104 Ile Glu Ser Leu Gly Ser Val Leu Phe Thr Gly Asn Leu Val Thr Pro 1045 1050 1055 Thr Leu Ser Thr Thr Thr Glu Gly Thr Pro Ala Thr Thr Ser Gly Asp 1060 1065 1070 Val Thr Lys Tyr Gly Ala Ala Ile Phe Gly Gln Ile Ala Ser Ser Asn 1075 1080 1085 Gly Ser Gln Thr Asp Asn Leu Pro Leu Lys Leu Ile Ala Ser Gly Gly 1090 1095 1100 Asn Ile Cys Phe Arg Asn Asn Glu Tyr Arg Pro Thr Ser Ser Asp Thr 1105 1110 1115 112 Gly Thr Ser Thr Phe Cys Ser Ile Ala Gly Asp Val Lys Leu Thr Met 1125 1130 1135 Gln Ala Ala Lys Gly Lys Thr Ile Ser Phe Phe Asp Ala Ile Arg Thr 1140 1145 1150Ser Thr Lys Lys Thr Gly Thr Gln Ala Thr Ala Tyr Asp Thr Leu Asp 1155 1160 1165 Ile Asn Lys Ser Glu Asp Ser Glu Thr Val Asn Ser Ala Phe Thr Gly 1170 1180 Thr Ile Leu Phe Ser Ser Glu Leu His Glu Asn Lys Ser Tyr Ile Pro 1185 1190 1195 1200 Gln Asn Val Val Leu His Ser Gly Ser Leu Val Leu Lys Pro Asn Thr 1205 1210 1215 Glu Leu His Val Ile Ser Phe Glu Gln Lys Glu Gly Ser Ser Leu Val 1220 1225 1230 Met Thr Pro Gly Ser Val Leu Ser Asn Gln Thr Val Ala Asp Gly Ala 1235 1240 1245 Leu Val Ile Asn Asn Met Thr Ile Asp Leu Ser Ser Val Glu Lys Asn 1250 1255 1260 Gly Ile Ala Glu Gly Asn Ile Phe Thr Pro Pro Glu Leu Arg Ile Ile 1265 1270 1275 1280 Asp Thr Thr Thr Ser Gly Ser Gly Gly Thr Pro Ser Thr Asp Ser Glu 1285 1290 1295 Ser Asn Gln Asn Ser Asp Asp Thr Lys Glu Gln Asn Asn Asn Asp Ala 1300 1305 1310 Ser Asn Gln Gly Glu Ser Ala Asn Gly Ser Ser Ser Pro Ala Val Ala 1315 1320 1325Ala Ala His Thr Ser Arg Thr Arg Asn Phe Ala Ala Ala Ala Thr Ala 1330 1335 1340 Thr Pro Thr Thr Thr Pro Thr Ala Thr Thr Thr Thr Ser Asn Gln Val 1345 1350 1355 Ile Leu Gly Gly Glu Ile Lys Leu Ile Asp Pro Asn Gly Thr Phe Phe

1365 1370 Gln Asn Fro Ala Leu Arg Ser Asp Gln Gln Ile Ser Leu Leu Val Leu 1380 1385 1390 Pro Thr Asp Ser Ser Lys Met Gln Ala Gln Lys Ile Val Leu Thr Gly 1395 1400 1405 Asp Ile Ala Pro Gln Lys Gly Tyr Thr Gly Thr Leu Thr Leu Asp Pro 1420 1410 1415 Asp Gln Leu Gln Asn Gly Thr Ile Ser Ala Leu Trp Lys Phe Asp Ser 1425 1430 1435 1440 Tyr Arg Gln Trp Ala Tyr Val Pro Arg Asp Asn His Phe Tyr Ala Asn 1445 1450 1450 Ser Ile Leu Gly Ser Gln Met Ser Met Val Thr Val Lys Gln Gly Leu 1460 1465 1470 Leu Asn Asp Lys Met Asn Leu Ala Arg Phe Asp Glu Val Ser Tyr Asn 1475 1480 1485 Asn Leu Trp Ile Ser Gly Leu Gly Thr Met Leu Ser Gln Val Gly Thr 1495 1500 1490 Pro Thr Ser Glu Glu Phe Thr Tyr Tyr Ser Arg Gly Ala Ser Val Ala 1505 1510 1515 1520 Leu Asp Ala Lys Pro Ala His Asp Val Ile Val Gly Ala Ala Phe Ser 1525 1530 1535 Lys Met Ile Gly Lys Thr Lys Ser Leu Lys Arg Glu Asn Asn Tyr Thr 1540 1545 1550 His Lys Gly Ser Glu Tyr Ser Tyr Gln Ala Ser Val Tyr Gly Gly Lys 1555 1560 1565 Pro Phe His Phe Val Ile Asn Lys Lys Thr Glu Lys Ser Leu Pro Leu 1575 1580 Leu Leu Gln Gly Val Ile Ser Tyr Gly Tyr Ile Lys His Asp Thr Val 1585 1590 1595 1600 Thr His Tyr Pro Thr Ile Arg Glu Arg Asn Gln Gly Glu Trp Glu Asp 1605 1610 1615 Leu Gly Trp Leu Thr Ala Leu Arg Val Ser Ser Val Leu Arg Thr Pro 1620 1625 1630 1620 Ala Gln Gly Asp Thr Lys Arg Ile Thr Val Tyr Gly Glu Leu Glu Tyr 1640 Ser Ser Ile Arg Gln Lys Gln Phe Thr Glu Thr Glu Tyr Asp Pro Arg 1650 1655. 1660 Tyr Phe Asp Asn Cys Thr Tyr Arg Asn Leu Ala Ile Pro Met Gly Leu 1665 1670 1675 168 Ala Phe Glu Gly Glu Leu Ser Gly Asn Asp Ile Leu Met Tyr Asn Arg 1695 1690 1695 Phe Ser Val Ala Tyr Met Pro Ser Ile Tyr Arg Asn Ser Pro Thr Cys 1700 1705 1710 Lys Tyr Gln Val Leu Ser Ser Gly Glu Gly Gly Glu Ile Ile Cys Gly 1715 1720 1725 Val Pro Thr Arg Asn Ser Ala Arg Gly Glu Tyr Ser Thr Gln Leu Tyr 1730 1735 . 1740 Pro Gly Pro Leu Trp Thr Leu Tyr Gly Ser Tyr Thr Ile Glu Ala Asp 1745 1750 1755 1760 Ala His Thr Leu Ala His Met Met Asn Cys Gly Ala Arg Met Thr Phe 1765 1770

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Thr Lou Phe Gln Glu Asn Thr Ala Lys Glu Glu Gly Gly Gly Leu Phe 520 Ile Lys Gly Thr Asp Lys Ala Leu Thr Met Thr Gly Leu Asp Ser Phe 535 Cys Leu Ile Asn Asn Thr Ser Glu Lys His Gly Gly Gly Ala Phe Val 545 550 555 560 Thr Lys Glu Ile Ser Gln Thr Tyr Thr Ser Asp Val Glu Thr Ile Pro Gly Ile Thr Pro Val His Gly Glu Thr Val Ile Thr Gly Asn Lys Ser 580 585 590 Thr Gly Gly Asn Gly Gly Gly Val Cys Thr Lys Arg Leu Ala Leu Ser 600 Asn Leu Gln Ser Ile Ser Ile Ser Gly Asn Ser Ala Ala Glu Asn Gly 610 615 620 Gly Gly Ala His Thr Cys Pro Asp Ser Phe Pro Thr Ala Asp Thr Ala 630 635 Glu Gln Pro Ala Ala Ala Ser Ala Ala Thr Ser Thr Pro Lys Ser Ala 645 650 Pro Val Ser Thr Ala Leu Ser Thr Pro Ser Ser Ser Thr Val Ser Ser 660 665 Leu Thr Leu Leu Ala Ala Ser Ser Gln Ala Ser Pro Ala Thr Ser Asn 675 680 685 Lys Glu Thr Gln Asp Pro Asn Ala Asp Thr Asp Leu Leu Ile Asp Tyr 695 Val Val Asp Thr Thr Ile Ser Lys Asn Thr Ala Lys Lys Gly Gly Gly 705 710 715 720 Ile Tyr Ala Lys Lys Ala Lys Met Ser Arg Ile Asp Gln Leu Asn Ile 725 730 735Ser Glu Asn Ser Ala Thr Glu Ile Gly Gly Gly Ile Cys Cys Lys Glu 740 745 750 Ser Leu Glu Leu Asp Ala Leu Val Ser Leu Ser Val Thr Glu Asn Leu 760 Val Gly Lys Glu Gly Gly Gly Leu His Ala Lys Thr Val Asn Ile Ser 770 775 780 Asn Leu Lys Ser Gly Phe Ser Phe Ser Asn Asn Lys Ala Asn Ser Ser 785 790 795 800 Ser Thr Gly Val Ala Thr Thr Ala Ser Ala Pro Ala Ala Ala Ala Ala 805 810 815 Ser Leu Gln Ala Ala Ala Ala Ala Ala Pro Ser Ser Pro Ala Thr Pro 820 825 830 Thr Tyr Ser Gly Val Val Gly Gly Ala Ile Tyr Gly Glu Lys Val Thr 835 . 840 845 Phe Ser Gln Cys Ser Gly Thr Cys Gln Phe Ser Gly Asn Gln Ala Ile 850 855 860 Asp Asn Asn Pro Ser Gln Ser Ser Leu Asn Val Gln Gly Gly Ala Ile 870 875 Tyr Ala Lys Thr Ser Leu Ser Ile Gly Ser Ser Asp Ala Gly Thr Ser 890 885 Tyr Ile Phe Ser Gly Asn Ser Val Ser Thr Gly Lys Ser Gln Thr Thr 905 900 910 Gly Gln Ile Ala Gly Gly Ala Ile Tyr Ser Pro Thr Val Thr Leu Asn 915 920 925 Cys Pro Ala Thr Phe Ser Asn Asn Thr Ala Ser Ile Ala Thr Pro Lys 930 935 940 Thr Ser Ser Glu Asp Gly Ser Ser Gly Asn Ser Ile Lys Asp Thr Ile 950 955 Gly Gly Ala Ile Ala Gly Thr Ala Ile Thr Leu Ser Gly Val Ser Arg 970 965 Phe Ser Gly Asn Thr Ala Asp Leu Gly Ala Ala Ile Gly Thr Leu Ala 985 Asn Ala Asn Thr Pro Ser Ala Thr Ser Gly Ser Gln Asn Ser Ile Thr

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Tyr Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Lys Pro Arg Gln Asp 1490 1495 1500 Phe Ile Leu Gly Ala Ala Phe Ser Lys Ile Val Gly Lys Thr Lys Ala 1510 1515 Ile Lys Lys Met His Asn Tyr Phe His Lys Gly Ser Glu Tyr Ser Tyr 1525 1530 1535 Gln Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Asn Lys 1540 1545 1550 Gln His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr 1555 1560 1565 Gly His Ile Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu 1570 1575 1580 Arg Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg 1585 1590 1595 1601 Ile Ser Met Asp Leu Lys Glu Pro Ser Lys Asp Ser Ser Lys Arg Ile 1605 1610 1615 Thr Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe 1620 1625 1630 Thr Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg 1635 1640 1645 Asn Leu Ser Leu Pro Val Gly Cys Ala Val Glu Gly Ala Ile Met Asn 1650 1660 Cys Asn Ile Leu Met Tyr Asn Lys Leu Ala Leu Ala Tyr Met Pro Ser 1665 1670 1685 Ile Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn 1685 1690 1695 Glu Ala Gly Gln Val Ile Cys Gly Val Pro Thr Arg Thr Ser Ala Arg 1700 1705 Ala Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr 1715 1720 1725 Gly Asn Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr 1735 Ser Cys Gly Ala Arg Met Ile Phe 1750 <210> 181 <211> 2601 <212> DNA <213> Chlamydia <400> 181 atggetagee ateaceatea ceateacete tttggecagg atecettagg tgaaacegee etecteacta aaaateetaa teatgtegte tgtacatttt ttgaggaetg taccatggag agcetettte etgetetttg tgeteatgea teacaagaeg atcetttgta tgtacttgga 180 aatteetaet gttqgttegt atetaaaete catateaeqq acceeaaaqa qqetettttt 240 aaagaaaaag gagatettte catteaaaae tttegettee ttteetteae agattgetet 300 tecaaggaaa geteteette tattatteat caaaagaatg gteagttate ettgegeaat 360 aatggtagea tgagtttetg tegaaateat getgaagget etggaggage eatetetgeg gatgeetttt etetacagea eaactatett tteacaget ttgaagagaa ttettetaaa 420 480 ggaaatggog gagecattea ggotcaaaco ttototttat otagaaatgt gtogootatt 540 tetttegece gtaategtge ggatttaaat ggeggegeta tttgetgtag taatettatt 600 tqttcaqqqa atqtaaaccc tctctttttc actqqaaact ccqccacraa tqqaqqcsct 660 attigtigta teagegatet aaacacetea gaaaaagget etetetetet tgettigtaac 720 780 caaraaacgc tatttgcaag caattctgct aaagaaaaag gcggggctat ttatgccaag cacatggtat tgcgttataa cggtcctgtt tccttcatta acaacagcgc taaaataggt 840 ggagctateg ceatecagte eggagggagt etetetatee ttgeaggtga aggatetgtt 900 ctgttccaga ataactccca acgcacctcc gaccaaggtc tagtaagaaa cgccatctac 960 ttagagaaag atgegattet ttetteetta gaagetegea acggagatat tetttettt 1020 gatectattg tacaagaaag tagcagcaaa gaategeete tteeeteete tttgcaagee agegtgactt cteccaccec agecaccqca teteetttag ttattcagac aagtgcaaac cgttcagtga ttttctcgag cgaacgtctt tctgaagaag aaaaaactcc tgataacctc 1200

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Trp Phe Val Ser Lys Leu His Ile Thr Asp Pro Lys Glu Ala Leu Phe
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Thr Asp Cys Ser Ser Lys Glu Ser Ser Pro Ser Ile Ile His Gln Lys
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Asn His Ala Glu Gly Ser Gly Gly Ala Ile Ser Ala Asp Ala Phe Ser
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Leu Gln His Asn Tyr Leu Phe Thr Ala Phe Glu Glu Asn Ser Ser Lys
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Phe Phe Thr Gly Asn Ser Ala Thr Asn Gly Gly Xaa Ile Cys Cys Ile
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Ser Asp Leu Asn Thr Ser Glu Lys Gly Ser Leu Ser Leu Ala Cys Asn
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Gln Xaa Thr Leu Phe Ala Ser Asn Ser Ala Lys Glu Lys Gly Gly Ala
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Phe 385	Ser	Ser	Glu	Arg	Leu 390	Ser	Glu	Glu	Glu	Lys 395	Thr	Pro	Asp	Asn	Leu 400
Thr	Ser	Gln	Leu	Gln 405	Gln	Pro	Ile	Glu	Leu 410	Lys	Ser	Gly	Arg	Leu 415	Val
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	Met			805					810					815	
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Gln Arg Ala Leu His Arg Leu Gln Thr Leu Leu Asn Val Ser Cys Val 835 840 845 Leu Arg Gly Gln Ser His Ser Tyr Ser Leu Asp Leu Gly Thr Thr Tyr 855 Arg Phe 865 <210> 190 <211> 1006 <212> PRT <213> Chlamydia <400> 190 Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Asp Ser Ser Leu 10 Val Pro His His His His His Met Ile Pro Gln Gly Ile Tyr Asp 20 Gly Glu Thr Leu Thr Val Ser Phe Pro Tyr Thr Val Ile Gly Asp Pro 40 Ser Gly Thr Thr Val Phe Ser Ala Gly Glu Leu Thr Leu Lys Asn Leu 55 Asp Asn Ser Ile Ala Ala Leu Pro Leu Ser Cys Phe Gly Asn Leu Leu 65 70 75 80 Gly Ser Phe Thr Val Leu Gly Arg Gly His Ser Leu Thr Phe Glu Asn 85 90 Ile Arg Thr Ser Thr Asn Gly Ala Ala Leu Ser Asn Ser Ala Ala Asp 105 110 Gly Leu Phe Thr Ile Glu Gly Phe Lys Glu Leu Ser Phe Ser Asn Cys 115 120 125 Asn Ser Leu Leu Ala Val Leu Pro Ala Ala Thr Thr Asn Lys Gly Ser 130 135 140 Gln Thr Pro Thr Thr Thr Ser Thr Pro Ser Asn Gly Thr Ile Tyr Ser 150 155 Lys Thr Asp Leu Leu Leu Asn Asn Glu Lys Phe Ser Phe Tyr Ser 165 170 Asn Leu Val Ser Gly Asp Gly Gly Ala Ile Asp Ala Lys Ser Leu Thr 180 185 190 Val Gln Gly Ile Ser Lys Leu Cys Val Phe Gln Glu Asn Thr Ala Gln 195 200 205 Ala Asp Gly Gly Ala Cys Gln Val Val Thr Ser Phe Ser Ala Met Ala 210 215 220 220 Asn Glu Ala Pro Ile Ala Phe Val Ala Asn Val Ala Gly Val Arg Gly 225 230 235 240 235 Gly Gly Ile Ala Ala Val Gln Asp Gly Gln Gln Gly Val Ser Ser Ser 245 250 255 Thr Ser Thr Glu Asp Pro Val Val Ser Phe Ser Arg Asn Thr Ala Val 265 Glu Phe Asp Gly Asn Val Ala Arg Val Gly Gly Gly Ile Tyr Ser Tyr 280 Gly Asn Val Ala Phe Leu Asn Asn Gly Lys Thr Leu Phe Leu Asn Asn 295 Val Ala Ser Pro Val Tyr Ile Ala Ala Lys Gln Pro Thr Ser Gly Gln 305 310 315 320 Ala Ser Asn Thr Ser Asn Asn Tyr Gly Asp Gly Gly Ala Ile Phe Cys 325 330 335 Lys Asn Gly Ala Gln Ala Gly Ser Asn Asn Ser Gly Ser Val Ser Phe 340 345 350350 Asp Gly Glu Gly Val Val Phe Phe Ser Ser Asn Val Ala Ala Gly Lys 360 365 Gly Gly Ala Ile Tyr Ala Lys Lys Leu Ser Val Ala Asn Cys Gly Pro

375 380

Val Gln Phs Leu Arg Asn Ile Ala Asn Asp Gly Gly Ala Ile Tyr Leu 390 395 Gly Glu Ser Gly Glu Leu Ser Leu Ser Ala Asp Tyr Gly Asp Ile Ile 405 410 Phe Asp Gly Asn Leu Lys Arg Thr Ala Lys Glu Asn Ala Ala Asp Val 420 425 430 Asn Gly Val Thr Val Ser Ser Gln Ala Ile Ser Met Gly Ser Gly Gly
435 440 445 Lys Ile Thr Thr Leu Arg Ala Lys Ala Gly His Gln Ile Leu Phe Asn 450 455 Asp Pro Ile Glu Met Ala Asn Gly Asn Asn Gln Pro Ala Gln Ser Ser 470 475 Lys Leu Leu Lys Ile Asn Asp Gly Glu Gly Tyr Thr Gly Asp Ile Val 490 485 Phe Ala Asn Gly Ser Ser Thr Leu Tyr Gln Asn Val Thr Ile Glu Gln 500 $$ 500 $$ Gly Arg Ile Val Leu Arg Glu Lys Ala Lys Leu Ser Val Asn Ser Leu 515 520 525Ser Gln Thr Gly Gly Ser Leu Tyr Met Glu Ala Gly Ser Thr Leu Asp 530 535 540 Phe Val Thr Pro Gln Pro Pro Gln Gln Pro Pro Ala Ala Asn Gln Leu 545 550 550 560 Ile Thr Leu Ser Asn Leu His Leu Ser Leu Ser Ser Leu Leu Ala Asn 565 570 575 Asn Ala Val Thr Asn Pro Pro Thr Asn Pro Pro Ala Gln Asp Ser His 580 585 590 Pro Ala Val Ile Gly Ser Thr Thr Ala Gly Ser Val Thr Ile Ser Gly 600 595 605 Pro Ile Phe Phe Glu Asp Leu Asp Asp Thr Ala Tyr Asp Arg Tyr Asp $610 \hspace{1.5cm} 620 \hspace{1.5cm} 620$ Trp Leu Gly Ser Asn Gln Lys Ile Asn Val Leu Lys Leu Gln Leu Gly 625 630 635 640 Thr Lys Pro Pro Ala Asn Ala Pro Ser Asp Leu Thr Leu Gly Asn Glu
645 650 655 Met Pro Lys Tyr Gly Tyr Gln Gly Ser Trp Lys Leu Ala Trp Asp Pro 660 665 670 Asn Thr Ala Asn Asn Gly Pro Tyr Thr Leu Lys Ala Thr Trp Thr Lys 675 680 685 Thr Gly Tyr Asn Pro Gly Pro Glu Arg Val Ala Ser Leu Val Pro Asn 690 695 700 Ser Leu Trp Gly Ser Ile Leu Asp Ile Arg Ser Ala His Ser Ala Ile 705 710 715 720 Gln Ala Ser Val Asp Gly Arg Ser Tyr Cys Arg Gly Leu Trp Val Ser 725 730 735 Gly Val Ser Asn Phe Phe Tyr His Asp Arg Asp Ala Leu Gly Gln Gly 740 745 750 Tyr Arg Tyr Ile Ser Gly Gly Tyr Ser Leu Gly Ala Asn Ser Tyr Phe 755 760 765 Gly Ser Ser Met Phe Gly Leu Ala Phe Thr Glu Val Phe Gly Arg Ser 770 780 Lys Asp Tyr Val Val Cys Arg Ser Asn His His Ala Cys Ile Gly Ser 785 790 795 Val Tyr Leu Ser Thr Gln Gln Ala Leu Cys Gly Ser Tyr Leu Phe Gly 805 810 815 Asp Ala Phe Ile Arg Ala Ser Tyr Gly Phe Gly Asn Gln His Met Lys 820 825 830 Thr Ser Tyr Thr Phe Ala Glu Glu Ser Asp Val Arg Trp Asp Asn Asn 835 840Cys Leu Ala Gly Glu Ile Gly Ala Gly Leu Pro Ile Val Ile Thr Pro 850 860 Ser Lys Leu Tyr Leu Asn Glu Leu Arg Pro Phe Val Gln Ala Glu Phe

<210> 191 <211> 977 <212> PRT

<213> Chlamydia

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Glu Ala Ser Asp Gly Gly Ala Ile Lys Val Thr Thr Arg Leu Asp Val

90

295 Thr Gly Asn Arg Gly Arg Ile Phe Phe Ser Asp Asn Ile Thr Lys Asn 305 310 315 Tyr Gly Gly Ala Ile Tyr Ala Pro Val Val Thr Leu Val Asp Asn Gly 325 330 335 Pro Thr Tyr Phe Ile Asn Asn Ile Ala Asn Asn Lys Gly Gly Ala Ile 340 345 350 Tyr Ile Asp Gly Thr Ser Asn Ser Lys Ile Ser Ala Asp Arg His Ala 355 360 365 Ile Ile Phe Asn Glu Asn Ile Val Thr Asn Val Thr Asn Ala Asn Gly 370 375 380 Thr Ser Thr Ser Ala Asn Pro Pro Arg Arg Asn Ala Ile Thr Val Ala 385 390 395 400 Ser Ser Ser Gly Glu Ile Leu Leu Gly Ala Gly Ser Ser Gln Asn Leu 405 410 415 Ile Phe Tyr Asp Pro Ile Glu Val Ser Asn Ala Gly Val Ser Val Ser 420 425 Phe Asn Lys Glu Ala Asp Gln Thr Gly Ser Val Val Phe Ser Gly Ala 440 Thr Val Asn Ser Ala Asp Phe His Gln Arg Asn Leu Gln Thr Lys Thr 450 455 460 Pro Ala Pro Leu Thr Leu Ser Asn Gly Phe Leu Cys Ile Glu Asp His 465 470 475 480 Ala Gln Leu Thr Val Asn Arg Phe Thr Gln Thr Gly Gly Val Val Ser 485 490 495 Leu Gly Asn Gly Ala Val Leu Ser Cys Tyr Lys Asn Gly Thr Gly Asp 500 505 510 Ser Ala Ser Asn Ala Ser Ile Thr Leu Lys His Ile Gly Leu Asn Leu 515 520 525 Ser Ser Ile Leu Lys Ser Gly Ala Glu Ile Pro Leu Trp Val Glu 530 535 Pro Thr Asn Asn Ser Asn Asn Tyr Thr Ala Asp Thr Ala Ala Thr Phe 550 555 Ser Leu Ser Asp Val Lys Leu Ser Leu Ile Asp Asp Tyr Gly Asn Ser 565 570 575 Pro Tyr Glu Ser Thr Asp Leu Thr His Ala Leu Ser Ser Gln Pro Met 580 585 590 Leu Ser Ile Ser Glu Ala Ser Asp Asn Gln Leu Gln Ser Glu Asn Ile 595 600 605 Asp Phe Ser Gly Leu Asn Val Pro His Tyr Gly Trp Gln Gly Leu Trp 610 620 Thr Trp Gly Trp Ala Lys Thr Gln Asp Pro Glu Pro Ala Ser Ser Ala 625 630 635 Thr Ile Thr Asp Pro Gln Lys Ala Asn Arg Phe His Arg Thr Leu Leu 645 650 655 Leu Thr Trp Leu Pro Ala Gly Tyr Val Pro Ser Pro Lys His Arg Ser 665 Pro Leu Ile Ala Asn Thr Leu Trp Gly Asn Met Leu Leu Ala Thr Glu 675 680 685 Ser Leu Lys Asn Ser Ala Glu Leu Thr Pro Ser Gly His Pro Phe Trp 690 695 700 Gly Ile Thr Gly Gly Gly Leu Gly Met Met Val Tyr Gln Asp Pro Arg 705 710 725 Glu Asn His Pro Gly Phe His Met Arg Ser Ser Gly Tyr Ser Ala Gly 725 730 735 Met Ile Ala Gly Gln Thr His Thr Phe Ser Leu Lys Phe Ser Gln Thr 740 745 750 Tyr Thr Lys Leu Asn Glu Arg Tyr Ala Lys Asn Asn Val Ser Ser Lys 755 760 765 Asn Tyr Ser Cys Gln Gly Glu Met Leu Phe Ser Leu Gln Glu Gly Phe
770 775 780

Leu Leu Thr Lys Leu Val Gly Leu Tyr Ser Tyr Gly Asp His Asn Cys 785 790 795 800 His His Phe Tyr Thr Gln Gly Glu Asn Leu Thr Ser Gln Gly Thr Phe 810 Arg Ser Gln Thr Met Gly Gly Ala Val Phe Phe Asp Leu Pro Met Lys 820 825 830Pro Phe Gly Ser Thr His Ile Leu Thr Ala Pro Phe Leu Gly Ala Leu Gly Ile Tyr Ser Ser Leu Ser His Phe Thr Glu Val Gly Ala Tyr Pro 855 860 Arg Ser Phe Ser Thr Lys Thr Pro Leu Ile Asn Val Leu Val Pro Ile 865 870 875 880 Gly Val Lys Gly Ser Phe Met Asn Ala Thr His Arg Pro Gln Ala Trp 885 890 895 Thr Val Glu Leu Ala Tyr Gln Pro Val Leu Tyr Arg Gln Glu Pro Gly 900 905 910 Ile Ala Thr Gln Leu Leu Ala Ser Lys Gly Ile Trp Phe Gly Ser Gly 915 920 925 Ser Pro Ser Ser Arg His Ala Met Ser Tyr Lys Ile Ser Gln Gln Thr 930 . 935 Gln Pro Leu Ser Trp Leu Thr Leu His Phe Gln Tyr His Gly Phe Tyr 945 950 955 960 Ser Ser Ser Thr Phe Cys Asn Tyr Leu Asn Gly Glu Ile Ala Leu Arg 965 970

<210> 192 <211> 848 <212> PRT <213> Chlamydia

<400> 192 Met Ala Ser His His His His His Gly Ala Ile Ser Cys Leu Arg Gly Asp Val Val Ile Ser Gly Asn Lys Gly Arg Val Glu Phe Lys Asp 20 25 30 Asn Ile Ala Thr Arg Leu Tyr Val Glu Glu Thr Val Glu Lys Val Glu 40 Glu Val Glu Pro Ala Pro Glu Gln Lys Asp Asn Asn Glu Leu Ser Phe 55 60 Leu Gly Ser Val Glu Gln Ser Phe Ile Thr Ala Ala Asn Gln Ala Leu 70 75 Phe Ala Ser Glu Asp Gly Asp Leu Ser Pro Glu Ser Ser Ile Ser Ser 85 90 95 85 Glu Glu Leu Ala Lys Arg Arg Glu Cys Ala Gly Gly Ala Ile Phe Ala 100 105 110 Lys Arg Val Arg Ile Val Asp Asn Gln Glu Ala Val Val Phe Ser Asn 120 125 Asn Phe Ser Asp Ile Tyr Gly Gly Ala Ile Phe Thr Gly Ser Leu Arg 130 135 Glu Glu Asp Lys Leu Asp Gly Gln Ile Pro Glu Val Leu Ile Ser Gly 145 150 155 160 Asn Ala Gly Asp Val Val Phe Ser Gly Asn Ser Ser Lys Arg Asp Glu 165 170 175 His Leu Pro His Thr Gly Gly Gly Ala Ile Cys Thr Gln Asn Leu Thr 180 185 Ile Ser Gln Asn Thr Gly Asn Val Leu Phe Tyr Asn Asn Val Ala Cys 195 200 205 Ser Gly Gly Ala Val Arg Ile Glu Asp His Gly Asn Val Leu Leu Glu 210 215 220

Ala Phe Gly Gly Asp Ile Val Phe Lys Gly Asn Ser Ser Phe Arg Ala 230 235 240 Gln Gly Ser Asp Ala Ile Tyr Phe Ala Gly Lys Glu Ser His Ile Thr 245 250 255 Ala Leu Asn Ala Thr Glu Gly His Ala Ile Val Phe His Asp Ala Leu 260 265 270 265 Val Phe Glu Asn Leu Lys Glu Arg Lys Ser Ala Glu Val Leu Leu Ile 275 280 285 Asn Ser Arg Glu Asn Pro Gly Tyr Thr Gly Ser Ile Arg Phe Leu Glu 295 Ala Glu Ser Lys Val Pro Gln Cys Ile His Val Gln Gln Gly Ser Leu 305 310 315 320 Glu Leu Leu Asn Gly Ala Thr Leu Cys Ser Tyr Gly Phe Lys Gln Asp \$325\$Ala Gly Ala Lys Leu Val Leu Ala Ala Gly Ser Lys Leu Lys Ile Leu 340 345 350 Asp Ser Gly Thr Pro Val Gln Gly His Ala Ile Ser Lys Pro Glu Ala 355 360 365 Glu Ile Glu Ser Ser Ser Glu Pro Glu Gly Ala His Ser Leu Trp Ile 370 375 380 Ala Lys Asn Ala Gln Thr Thr Val Pro Met Val Asp Ile His Thr Ile 385 390 395 400 Ser Val Asp Leu Ala Ser Phe Ser Ser Ser Gln Gln Glu Gly Thr Val 405 410 415Glu Ala Pro Gln Val Ile Val Pro Gly Gly Ser Tyr Val Arg Ser Gly
420 425 430 Glu Leu Asn Leu Glu Leu Val Asn Thr Thr Gly Thr Gly Tyr Glu Asn 435 440 445 His Ala Leu Leu Lys Asn Glu Ala Lys Val Pro Leu Met Ser Phe Val 455 Ala Ser Ser Asp Glu Ala Ser Ala Glu Ile Ser Asn Leu Ser Val Ser 465 470 475 480 Asp Leu Gln Ile His Val Ala Thr Pro Glu Ile Glu Glu Asp Thr Tyr 485 490 495 Gly His Met Gly Asp Trp Ser Glu Ala Lys Ile Gln Asp Gly Thr Leu 500 505 510 Val Ile Asn Trp Asn Pro Thr Gly Tyr Arg Leu Asp Pro Gln Lys Ala 515 520 525 Gly Ala Leu Val Phe Asn Ala Leu Trp Glu Glu Gly Ala Val Leu Ser 530 540 Ala Leu Lys Asn Ala Arg Phe Ala His Asn Leu Thr Ala Gln Arg Met 545 550 555 560Glu Phe Asp Tyr Ser Thr Asn Val Trp Gly Phe Ala Phe Gly Gly Phe
565 570 575 Arg Thr Leu Ser Ala Glu Asn Leu Val Ala Ile Asp Gly Tyr Lys Gly 580 585 590 Ala Tyr Gly Gly Ala Ser Ala Gly Val Asp Ile Gln Leu Met Glu Asp 595 600 605 Phe Val Leu Gly Val Ser Gly Ala Ala Phe Leu Gly Lys Met Asp Ser 610 620 Gln Lys Phe Asp Ala Glu Val Ser Arg Lys Gly Val Val Gly Ser Val 625 630 635 640 Tyr Thr Gly Phe Leu Ala Gly Ser Trp Phe Phe Lys Gly Gln Tyr Ser 645 650 655 Leu Gly Glu Thr Gln Asn Asp Met Lys Thr Arg Tyr Gly Val Leu Gly 665 Glu Ser Ser Ala Ser Trp Thr Ser Arg Gly Val Leu Ala Asp Ala Leu 675 680 685 Val Glu Tyr Arg Ser Leu Val Gly Pro Val Arg Pro Thr Phe Tyr Ala 690 700 Leu His Phe Asn Pro Tyr Val Glu Val Ser Tyr Ala Ser Met Lys Phe

705 | 710 | 710 | 715 | 720 | 725 | 725 | 726 | 726 | 726 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 | 727 |

<210> 193 <211> 778 <212> PRT <213> Chlamydia

<400> 193 Met His His His His His Gly Leu Ala Ser Cys Val Asp Leu His 10 Ala Gly Gly Gln Ser Val Asn Glu Leu Val Tyr Val Gly Pro Gln Ala 20 25 Val Leu Leu Asp Gln Ile Arg Asp Leu Phe Val Gly Ser Lys Asp 40 Ser Gin Ala Glu Gly Gin Tyr Arg Leu Ile Val Gly Asp Pro Ser Ser 55 Phe Gln Glu Lys Asp Ala Asp Thr Leu Pro Gly Lys Val Glu Gln Ser 65 70 75 80Thr Leu Phe Ser Val Thr Asn Pro Val Val Phe Gln Gly Val Asp Gln Gln Asp Gln Val Ser Ser Gln Gly Leu Ile Cys Ser Phe Thr Ser Ser 100 105 110Asn Leu Asp Ser Pro Arg Asp Gly Glu Ser Phe Leu Gly Tle Ala Phe 115 120 125 Val Gly Asp Ser Ser Lys Ala Gly Ile Thr Leu Thr Asp Val Lys Ala 130 135 140 Ser Leu Ser Gly Ala Ala Leu Tyr Ser Thr Glu Asp Leu Ile Phe Glu 150 155 Lys Ile Lys Gly Gly Leu Glu Phe Ala Ser Cys Ser Ser Leu Glu Gln 165 170 175 Gly Gly Ala Cys Ala Ala Gln Ser Ile Leu Ile His Asp Cys Gln Gly 180 185 190 Leu Gln Val Lys His Cys Thr Thr Ala Val Asn Ala Glu Gly Ser Ser 195 200 205 Ala Asn Asp His Leu Gly Phe Gly Gly Gly Ala Phe Phe Val Thr Gly 210 215 Ser Leu Ser Gly Glu Lys Ser Leu Tyr Met Pro Ala Gly Asp Met Val 225 230 240 Val Ala Asn Cys Asp Gly Ala Ile Ser Phe Glu Gly Asn Ser Ala Asn 245 250 Phe Ala Asn Gly Gly Ala Ile Ala Ala Ser Gly Lys Val Leu Phe Val 260 265 270 Ala Asn Asp Lys Lys Thr Ser Phe Ile Glu Asn Arg Ala Leu Ser Gly 275 280 285 Gly Ala Ile Ala Ala Ser Ser Asp Ile Ala Phe Gln Asn Cys Ala Glu

295 Leu Val Phe Lys Gly Asn Cys Ala Ile Gly Thr Glu Asp Lys Gly Ser 305 310 315 320Leu Gly Gly Gly Ala Ile Ser Ser Leu Gly Thr Val Leu Leu Gln Gly 325 330 335 Asn His Gly Ile Thr Cys Asp Lys Asn Glu Ser Ala Ser Gln Gly Gly 340 345 350 Ala Ile Phe Gly Lys Asn Cys Gln Ile Ser Asp Asn Glu Gly Pro Val 355 360 365 Val Phe Arg Asp Ser Thr Ala Cys Leu Gly Gly Gly Ala Ile Ala Ala 370 375 380 Gln Glu Ile Val Ser Ile Gln Asn Asn Gln Ala Gly Ile Ser Phe Glu 385 390 395 400 Gly Gly Lys Ala Ser Phe Gly Gly Gly Ile Ala Cys Gly Ser Phe Ser 405 410 415 Ser Ala Gly Gly Ala Ser Val Leu Gly Thr Ile Asp Ile Ser Lys Asn 420 425 430 Leu Gly Ala Ile Ser Phe Ser Arg Thr Leu Cys Thr Thr Ser Asp Leu 440 Ser Leu Ser Glu Asn Ala Gly Val Leu Thr Phe Lys Asp Asn Ile Val 465 470 480 Lys Thr Phe Ala Ser Asn Gly Lys Ile Leu Gly Gly Gly Ala Ile Leu 490 Ala Thr Gly Lys Val Glu Ile Thr Asn Asn Ser Gly Gly Ile Ser Phe $500 \hspace{1.5cm} 505 \hspace{1.5cm} 510$ Thr Gly Asn Ala Arg Ala Pro Gln Ala Leu Pro Thr Gln Glu Glu Phe 515 520 525Pro Leu Phe Ser Lys Lys Glu Gly Arg Pro Leu Ser Ser Gly Tyr Ser 530 540 Gly Gly Gly Ala Ile Leu Gly Arg Glu Val Ala Ile Leu His Asn Ala 545 550 560 Ala Val Val Phe Glu Gln Asn Arg Leu Gln Cys Ser Glu Glu Glu Ala 565 570 575 Thr Leu Leu Gly Cys Cys Gly Gly Gly Ala Val His Gly Met Asp Ser 580 585 590 Thr Ser Ile Val Gly Asn Ser Ser Val Arg Phe Gly Asn Asn Tyr Ala 595 600 605 Met Gly Gln Gly Val Ser Gly Gly Ala Leu Leu Ser Lys Thr Val Gln 610 615 620 Leu Ala Gly Asn Gly Ser Val Asp Phe Ser Arg Asn Ile Ala Ser Leu 625 630 635 640 Gly Gly Gly Ala Leu Gln Ala Ser Glu Gly Asn Cys Glu Leu Val Asp
645 655 Asn Gly Tyr Val Leu Phe Arg Asp Asn Arg Gly Arg Val Tyr Gly Gly 665 Ala Ile Ser Cys Leu Arg Gly Asp Val Val Ile Ser Gly Asn Lys Gly 675 680 685 Arg Val Glu Phe Lys Asp Asn Ile Ala Thr Arg Leu Tyr Val Glu Glu
690 695 700 Thr Val Glu Lys Val Glu Glu Val Glu Pro Ala Pro Glu Gln Lys Asp 705 710 715 720 Asn Asn Glu Leu Ser Phe Leu Gly Ser Val Glu Gln Ser Phe Ile Thr 725 730 735 Ala Ala As
n Gl
n Ala Leu Phe Ala Ser Glu Asp Gly Asp Leu Ser Pro $740 \hspace{1.5cm} 745 \hspace{1.5cm} 750 \hspace{1.5cm} 750$ Glu Ser Ser Ile Ser Ser Glu Glu Leu Ala Lys Arg Arg Glu Cys Ala 760 Gly Gly Ala Asp Ser Ser Arg Ser Gly Cys 770 775

<210> 194 <211> 948 <212> PRT <213> Chlamydia

<400> 194

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Val Glu Lys Asn Gly Ile Ala Glu Gly Asn Ile Phe Thr Pro Pro Glu 435 440 Leu Arg Ile Ile Asp Thr Thr Thr Ser Gly Ser Gly Gly Thr Pro Ser 450 460 Thr Asp Ser Glu Ser Asn Gln Asn Ser Asp Asp Thr Lys Glu Gln Asn 470 475 Asn Asn Asp Ala Ser Asn Gln Gly Glu Ser Ala Asn Gly Ser Ser Ser 485 490 495 Pro Ala Val Ala Ala Ala His Thr Ser Arg Thr Arg Asn Phe Ala Ala 500 505 510 Ala Ala Thr Ala Thr Pro Thr Thr Thr Pro Thr Ala Thr Thr Thr Thr 515 520 525 Ser Asn Gln Val Ile Leu Gly Gly Glu Ile Lys Leu Ile Asp Pro Asn 530 540 Gly Thr Phe Phe Gln Asn Pro Ala Leu Arg Ser Asp Gln Gln Ile Ser 550 555 Leu Leu Val Leu Pro Thr Asp Ser Ser Lys Met Gln Ala Gln Lys Ile 565 570 575 Val Leu Thr Gly Asp Ile Ala Pro Gln Lys Gly Tyr Thr Gly Thr Leu 580 585 590 Thr Leu Asp Pro Asp Gln Leu Gln Asn Gly Thr Ile Ser Ala Leu Trp 595 600 605 Lys Phe Asp Ser Tyr Arg Gln Trp Ala Tyr Val Pro Arg Asp Asn His 610 620 Phe Tyr Ala Asn Ser Ile Leu Gly Ser Gln Met Ser Met Val Thr Val 625 630 635 Lys Gln Gly Leu Leu Asn Asp Lys Met Asn Leu Ala Arg Phe Asp Glu 645 650 655 Val Ser Tyr Asn Asn Leu Trp Ile Ser Gly Leu Gly Thr Met Leu Ser 665 Gln Val Gly Thr Pro Thr Ser Glu Glu Phe Thr Tyr Tyr Ser Arg Gly 675 680 685 Ala Ser Val Ala Leu Asp Ala Lys Pro Ala His Asp Val Ile Val Gly 690 695 700 Ala Ala Phe Ser Lys Met Ile Gly Lys Thr Lys Ser Leu Lys Arg Glu 705 710 715 720 Asn Asn Tyr Thr His Lys Gly Ser Glu Tyr Ser Tyr Gln Ala Ser Val 725 730 735 Tyr Gly Gly Lys Pro Phe His Phe Val Ile Asn Lys Lys Thr Glu Lys 740 745 750 Ser Leu Pro Leu Leu Gln Gly Val Ile Ser Tyr Gly Tyr Ile Lys 755 760 765 His Asp Thr Val Thr His Tyr Pro Thr Ile Arg Glu Arg Asn Gln Gly 770 775 780 Glu Trp Glu Asp Leu Gly Trp Leu Thr Ala Leu Arg Val Ser Ser Val 785 790 795 800 Leu Arg Thr Pro Ala Gln Gly Asp Thr Lys Arg Ile Thr Val Tyr Gly 805 810 Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe Thr Glu Thr Glu 820 825 Tyr Asp Pro Arg Tyr Phe Asp Asn Cys Thr Tyr Arg Asn Leu Ala Ile 835 840 845 Pro Met Gly Leu Ala Phe Glu Gly Glu Leu Ser Gly Asn Asp Ile Leu 850 855 860 Met Tyr Asn Arg Phe Ser Val Ala Tyr Met Pro Ser Ile Tyr Arg Asn 865 870 875 880 Ser Pro Thr Cys Lys Tyr Gln Val Leu Ser Ser Gly Glu Gly Gly Glu 885 890 895 Ile Ile Cys Gly Val Pro Thr Arg Asn Ser Ala Arg Gly Glu Tyr Ser 900 905 910 Thr Gln Leu Tyr Pro Gly Pro Leu Trp Thr Leu Tyr Gly Ser Tyr Thr

920 Ile Glu Ala Asp Ala His Thr Leu Ala His Met Met Asn Cys Gly Ala 930 935 Arq Met Thr Phe 945 <210> 195 <211> 821 <212> PRT <213> Chlamydia <400> 195 Met His His His His His Glu Ala Ser Ser Ile Gln Asp Gln Ile 10 Lys Asn Thr Asp Cys Asn Val Ser Lys Val Gly Tyr Ser Thr Ser Gln 20 Ala Phe Thr Asp Met Met Leu Ala Asp Asn Thr Glu Tyr Arg Ala Ala 40 Asp Ser Val Ser Phe Tyr Asp Phe Ser Thr Ser Ser Gly Leu Pro Arg 55 Lys His Leu Ser Ser Ser Ser Glu Ala Ser Pro Thr Thr Glu Gly Val 65 70 75 80Ser Ser Ser Ser Ser Gly Glu Asn Thr Glu Asn Ser Gln Asp Ser Ala 85 90 95 Pro Ser Ser Gly Glu Thr Asp Lys Lys Thr Glu Glu Glu Leu Asp Asn 100 105 110Gly Gly Ile Ile Tyr Ala Arg Glu Lys Leu Thr Ile Ser Glu Ser Gln 120 Asp Ser Leu Ser Asn Pro Ser Ile Glu Leu His Asp Asn Ser Phe Phe 135 140 Phe Gly Glu Gly Glu Val Ile Phe Asp His Arg Val Ala Leu Lys Asn 155 150 Gly Gly Ala Ile Tyr Gly Glu Lys Glu Val Val Phe Glu Asn Ile Lys 165 170 175 Ser Leu Leu Val Glu Val Asn Ile Ser Val Glu Lys Gly Gly Ser Val Tyr Ala Lys Glu Arg Val Ser Leu Glu Asn Val Thr Glu Ala Thr Phe 195 200 205 Ser Ser Asn Gly Gly Glu Gln Gly Gly Gly Gly Ile Tyr Ser Glu Gln 215 Asp Met Leu Ile Ser Asp Cys Asn Asn Val His Phe Gln Gly Asn Ala 225 230 235 240 Ala Gly Ala Thr Ala Val Lys Gln Cys Leu Asp Glu Glu Met Ile Val 245 250 Leu Leu Thr Glu Cys Val Asp Ser Leu Ser Glu Asp Thr Leu Asp Ser 260 270 265 Thr Pro Glu Thr Glu Gln Thr Lys Ser Asn Gly Asn Gln Asp Gly Ser 280 Ser Glu Thr Lys Asp Thr Gln Val Ser Glu Ser Pro Glu Ser Thr Pro 295 300 Ser Pro Asp Asp Val Leu Gly Lys Gly Gly Gly Ile Tyr Thr Glu Lys Ser Leu Thr Ile Thr Gly Ile Thr Gly Thr Ile Asp Phe Val Ser Asn 325 330 335 Ile Ala Thr Asp Ser Gly Ala Gly Val Phe Thr Lys Glu Asn Leu Ser 340 345 350 Cys Thr Asn Thr Asn Ser Leu Gln Phe Leu Lys Asn Ser Ala Gly Gln
355 360 365 His Gly Gly Gly Ala Tyr Val Thr Gln Thr Met Ser Val Thr Asn Thr 370 375 380 375 Thr Ser Glu Ser Ile Thr Thr Pro Pro Leu Val Gly Glu Val Ile Phe

385					390					395					400
	Glu	Asn	Thr	Ala 405		Gly	His	Gly	Gly 410		Ile	Cys	Thr	Asn 415	
Leu	Ser	Leu	Ser 420		Leu	Lys	Thr	Val 425		Leu	Thr	Lys	Asn 430		Ala
Lys	Glu	Ser 435	Gly	Gly	Ala	Ile	Phe 440	Thr	Asp	Leu	Ala	Ser	Ile	Pro	Thr
Thr	Asp 450	Thr	Pro	Glu	Ser	Ser 455	Thr	Pro	Ser	Ser	Ser 460	Ser	Pro	Ala	Ser
465			Val		470			-		475	-				480
			Pro	485					490					495	
			Thr 500					505			-		510		
		515	Ile				520					525			-
-	530	_	Ala		-	535		_		-	540		_		
545			Leu		550					555		_		_	560
			Glu	565					570					575	
	_		Ser 580			-		585	-				590	_	
		595	Ser			_	600					605	-		
	610		Ile			615					620				
625			Gly		630					635					640
		-	Ser	645					650		-			655	-
		-	Thr 660	-		-		665					670		
		675	Gly				680	-				685	-		
	690		Glu			695					700		_		
705			Thr	_	710			-		715					720
_			Val Ala	725				-	730	-				735	
_	_	-	740				-	745			-	_	750		
		755	Ala				760					765			
	770		Ser			775					780				
785			Ser		790					795					800
			Pro	805	нта	GTÄ	ser	ınr	810	GIU	ınr	FEO	THE	815	тте
оту	сту	оту	Ala 820	TTE											

<210> 196 <211> 525 <212> PRT <213> Chlamydia

<400> 196 Met His His His His His Thr Ala Ala Ser Asp Asn Phe Gln Leu Ser Gln Gly Gly Gln Gly Phe Ala Ile Pro Ile Gly Gln Ala Met Ala 25 Ile Ala Gly Gln Ile Lys Leu Pro Thr Val His Ile Gly Pro Thr Ala 40 4.5 Phe Leu Gly Leu Gly Val Val Asp Asn Asn Gly Asn Gly Ala Arg Val 55 Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr 70 75 Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr 90 85 Ala Met Ala Asp Ala Leu Asn Gly His His Pro Gly Asp Val Ile Ser 100 105 Val Thr Trp Gln Thr Lys Ser Gly Gly Thr Arg Thr Gly Asn Val Thr 115 120 Leu Ala Glu Gly Pro Pro Ala Glu Phe Pro Leu Val Pro Arg Gly Ser 135 140 Pro Leu Pro Val Gly Asn Pro Ala Glu Pro Ser Leu Leu Ile Asp Gly 145 150 155 Thr Met Trp Glu Gly Ala Ser Gly Asp Pro Cys Asp Pro Cys Ala Thr 165 170 Trp Cys Asp Ala Ile Ser Ile Arg Ala Gly Tyr Tyr Gly Asp Tyr Val 180 185 190 Phe Asp Arg Val Leu Lys Val Asp Val Asn Lys Thr Phe Ser Gly Met 200 Ala Ala Thr Pro Thr Gln Ala Ile Gly Asn Ala Ser Asn Thr Asn Gln 215 220 Pro Glu Ala Asn Gly Arg Pro Asn Ile Ala Tyr Gly Arg His Met Gln 230 235 Asp Ala Glu Trp Phe Ser Asn Ala Ala Phe Leu Ala Leu Asn Ile Trp 245 250 255Asp Arg Phe Asp Ile Phe Cys Thr Leu Gly Ala Ser Asn Gly Tyr Phe 265 Lys Ala Ser Ser Ala Ala Phe Asn Leu Val Gly Leu Ile Gly Phe Ser 275 280 285 Ala Ala Ser Ser Ile Ser Thr Asp Leu Pro Met Gln Leu Pro Asn Val Gly Ile Thr Gln Gly Val Val Glu Phe Tyr Thr Asp Thr Ser Phe Ser 310 315 Trp Ser Val Gly Ala Arg Gly Ala Leu Trp Glu Cys Gly Cys Ala Thr 325 330 335 Leu Gly Ala Glu Phe Gln Tyr Ala Gln Ser Asn Pro Lys Ile Glu Met 340 345 350 Leu Asn Val Thr Ser Ser Pro Ala Gln Phe Val Ile His Lys Pro Arg 355 360 Gly Tyr Lys Gly Ala Ser Ser Asn Phe Pro Leu Pro Ile Thr Ala Gly 370 375 380Thr Thr Glu Ala Thr Asp Thr Lys Ser Ala Thr Ile Lys Tyr His Glu 390 Trp Gln Val Gly Leu Ala Leu Ser Tyr Arg Leu Asn Met Leu Val Pro 405 410 415 Tyr Ile Gly Val Asn Trp Ser Arg Ala Thr Phe Asp Ala Asp Thr Ile 420 425 430 Arg Ile Ala Gln Pro Lys Leu Lys Ser Glu Ile Leu Asn Ile Thr Thr 440 Trp Asn Pro Ser Leu Ile Gly Ser Thr Thr Ala Leu Pro Asn Asn Ser 455 460 Gly Lys Asp Val Leu Ser Asp Val Leu Gln Ile Ala Ser Ile Gln Ile

Asn Lys Met Lys Ser Arg Lys Ala Cys Gly Val Ala Val Gly Ala Thr 485 490 495	
Leu Ile Asp Ala Asp Lys Trp Ser Ile Thr Gly Glu Ala Arg Leu Ile 500 505 510	
Asn Glu Arg Ala Ala His Met Asn Ala Gln Phe Arg Phe 515 520 525	
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<400> 197 gataggogog cogoaatcat gaaatttatg toagctactg otg	43
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<210> 199 <211> 6 <212> DNA <213> Chlamydia	
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<210> 200 <211> 34 <212> DNA <213> Chlamydia	
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<210> 202 <211> 5 <212> DNA <213> Chlamydia	
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tgcaatcatg	aaaaaagcgt	ttttctttt	С			31
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<213> Chlamydia	
<400> 211 cagacatatg catcaccatc accatcacgg gttagc	36
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<210> 213 <211> 51 <212> DNA <213> Chlamydia	
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<210> 216 <211> 31 <212> DNA <213> Chlamydia	
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Val Pro Ser Ser Asp Pro
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<210> 219
<211> 51
<212> DNA
<213> Chlamydia
<400> 219
cagaggtacc gcatcaccat caccatcaca tgattcctca aggaatttac g
                                                                       51
<210> 220
<211> 33
<212> DNA
<213> Chlamydia
<400> 220
cagageggee gettagaace ggaetttact tee
                                                                        33
<210> 221
<211> 24
<212> PRT
<213> Chlamydia
<400> 221
Met Ala Ser Met Thr Gly Gly Gln Gln Asn Gly Arg Asp Ser Ser Leu
                 5
                                    10
Val Pro His His His His His His
            20
<210> 222
<211> 46
<212> DNA
<213> Chlamydia
<400> 222
cagagetage cateaceate accateacet etttggecag gatece
                                                                        46
<210> 223
<211> 30
<212> DNA
<213> Chlamydia
<400> 223
cagaactagt ctagaacctg taagtggtcc
                                                                        30
<210> 224
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 224
Met Ser Gln Lys Asn Lys Asn Ser Ala Phe Met His Pro Val Asn Ile
                 5
1
                                    10
                                                        15
Ser Thr Asp Leu
            20
<210> 225
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<211> 20
<212> PRT
<213> Artificial Sequence
<223> Made in a lab
<400> 225
Lys Asn Ser Ala Phe Met His Pro Val Asn Ile Ser Thr Asp Len Ala
                                    10
Val Ile Val Glv
            20
<210> 226
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 226
His Pro Val Asn Ile Ser Thr Asp Leu Ala Val Ile Val Gly Lys Gly
1
                                    10
Pro Met Pro Arg
            20
<210> 227
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 227
Ser Thr Asp Leu Ala Val Ile Val Gly Lys Gly Pro Met Pro Arg Thr
1
Glu Ile Val Lys
            20
<210> 228
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 228
Val Ile Val Gly Lys Gly Pro Met Pro Arg Thr Glu Ile Val Lys Lys
Val Trp Glu Tyr
<210> 229
<211> 20
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<213> Artificial Sequence
<220>
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<223> Made in a lab
<400> 229
Gly Pro Met Pro Arg Thr Glu Ile Val Lys Lys Val Trp Glu Tyr Ile
1
Lys Lys His Asn
           20
<210> 230
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 230
Ile Lys Lys His Asn Cys Gln Asp Gln Lys Asn Lys Arg Asn Ile Leu
                                  10
Pro Asp Ala Asn
            20
<210> 231
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 231
Asn Cys Gln Asp Gln Lys Asn Lys Arg Asn Ile Leu Pro Asp Ala Asn
                                   10
Leu Ala Lys Val
<210> 232
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 232
Lys Asn Lys Arg Asn Ile Leu Pro Asp Ala Asn Leu Ala Lys Val Phe
Gly Ser Ser Asp
            20
<210> 233
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 233
Ile Leu Pro Asp Ala Asn Leu Ala Lys Val Phe Gly Ser Ser Asp Pro
1 5
                                  10
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Ile Asp Met Phe
<210> 234
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 234
Asn Leu Ala Lys Val Phe Gly Ser Ser Asp Pro Ile Asp Met Phe Gln
Met Thr Lys Ala
            20
<210> 235
<211> 22
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 235
Phe Gly Ser Ser Asp Pro Ile Asp Met Phe Gln Met Thr Lys Ala Leu
1
                5
                                   10
Ser Lys His Ile Val Lys
<210> 236
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 236
Val Glu Ile Thr Gln Ala Val Pro Lys Tyr Ala Thr Val Gly Ser Pro
                                   10
Tyr Pro Val Glu
            20
<210> 237
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 237
Ala Val Pro Lys Tyr Ala Thr Val Gly Ser Pro Tyr Pro Val Glu Ile
1
Thr Ala Thr Gly
<210> 238
<211> 20
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<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 238
Ala Thr Val Gly Ser Pro Tyr Pro Val Glu Ile Thr Ala Thr Gly Lys
                                    10
Arg Asp Cys Val
<210> 239
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 239
Pro Tyr Pro Val Glu Ile Thr Ala Thr Gly Lys Arg Asp Cys Val Asp
                                   10
Val Ile Ile Thr
            20
<210> 240
<211> 21
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 240
Ile Thr Ala Thr Gly Lys Arg Asp Cys Val Asp Val Ile Ile Thr Gln
1
                                    10
Gln. Leu Pro Cys Glu
<210> 241
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 241
Lys Arg Asp Cys Val Asp Val Ile Ile Thr Gln Gln Leu Pro Cys Glu
 ī
Ala Glu Phe Val
<210> 242
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
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<400> 242
Asp Val Ile Ile Thr Gln Gln Leu Pro Cys Glu Ala Glu Phe Val Arg
Ser Asp Pro Ala
           20
<210> 243
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 243
Thr Gln Gln Leu Pro Cys Glu Ala Glu Phe Val Arg Ser Asp Pro Ala
                                    10
Thr Thr Pro Thr
          20
<210> 244
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 244
Cys Glu Ala Glu Phe Val Arg Ser Asp Pro Ala Thr Thr Pro Thr Ala
                                    10
Asp Gly Lys Leu
           20
<210> 245
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 245
Val Arg Ser Asp Pro Ala Thr Thr Pro Thr Ala Asp Gly Lys Leu Val
                                    10
Trp Lys Ile Asp
            20
<210> 246
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 246
Ala Thr Thr Pro Thr Ala Asp Gly Lys Leu Val Trp Lys Ile Asp Arg
                                    10
Leu Gly Gln Gly
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<212> PRT

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<210> 247
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 247
Ala Asp Gly Lys Leu Val Trp Lys Ile Asp Arg Leu Gly Gln Gly Glu
Lys Ser Lys Ile
            20
<210> 248
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 248
Val Trp Lys Ile Asp Arg Leu Gly Gln Gly Glu Lys Ser Lys Ile Thr
                                    10
Val Trp Val Lys
            20
<210> 249
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 249
Arg Leu Gly Gln Gly Glu Lys Ser Lys Ile Thr Val Trp Val Lys Pro
1
                                    10
Leu Lys Glu Gly
<210> 250
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 250
Gly Glu Lys Ser Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
1
                                   10
Cys Cys Phe Thr
            20
<210> 251
<211> 16
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<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 251
Gly Glu Lys Ser Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
<210> 252
<211> 12
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 252
Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
<210> 253
<211> 16
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 253
Gly Asp Lys Cys Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly
<210> 254
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 254
Thr Glu Tyr Pro Leu Leu Ala Asp Pro Ser Phe Lys Ile Ser Glu Ala
                 5
                                    10
Phe Gly Val Leu
            20
<210> 255
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 255
Leu Ala Asp Pro Ser Phe Lys Ile Ser Glu Ala Phe Gly Val Leu Asn
                                   10
Pro Glu Gly Ser
            20
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<210> 256
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 256
Phe Lys Ile Ser Glu Ala Phe Gly Val Leu Asn Pro Glu Gly Ser Leu
Ala Leu Arg Ala
            20
<210> 257
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 257
Ala Phe Gly Val Leu Asn Pro Glu Gly Ser Leu Ala Leu Arg Ala Thr
1
                                   10
                                                       1.5
Phe Leu Ile Asp
            20
<210> 258
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 258
Asn Pro Glu Gly Ser Leu Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys
                                   10
His Gly Val Ile
<210> 259
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 259
Leu Ala Leu Arg Ala Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg
1
                 5
                                  10
His Ala Val Ile
           20
<210> 260
<211> 20
<212> PRT
<213> Artificial Sequence
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<220>
<223> Made in a lab
<400> 260
Thr Phe Leu Ile Asp Lys His Gly Val Ile Arg His Ala Val Ile Asp
                                     10
Asp Leu Pro Leu
            20
<210> 261
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 261
Lys His Gly Val Ile Arg His Ala Val Ile Asn Asp Leu Pro Leu Gly
                                    10
Arg Ser Ile Asp
            20
<210> 262
<211> 20
<212> PRT
<213> Artificial Sequence
<220>
<223> Made in a lab
<400> 262
Arg His Ala Val Ile Asn Asp Leu Pro Leu Gly Arg Ser Ile Asp Glu
                                    10
Glu Leu Arg Ile
<210> 263
<211> 897
<212> DNA
<213> Chlamydia
<220>
<221> misc feature
<222> 604
<223> n = A,T,C or G
<400> 263
atggetteta tatgeggacg tttagggtet ggtacaggga atgetetaaa agettttttt
acacaqccca acaataaaat ggcaagggta gtaaataaga cgaagggagt ggataagact
                                                                         120
attaaggttg ccaagtctgc tgccgaattg accgcaaata ttttggaaca agctggaggc
                                                                        180
gegggetett cegeacacat tacagettee caaqtqteea aaqqattagg ggatqegaga
                                                                        240
actifttgteg etttagggaa tgeetttaac ggagegttge caggaacagt teaaagtgeg
                                                                        300
casagettet teteteacat gaaagetget agteagaaaa egcaagaagg ggatgagggg
                                                                        360
ctcacagcag atctttgtgt gtctcataag cgcagagcgg ctgcggctgt ctgtagcatc
                                                                        420
atoggaggaa ttacctacct cgcgacattc ggagctatcc gtccgattct gtttgtcaac aaaatgctgg caaaaccgtt tctttcttcc caaactaaag caaaatggg atcttctgtt
                                                                        480
                                                                        540
agctatatta tggcggctaa ccatgcageg tetgtggtgg gtgctggact cgctateagt
                                                                        600
gegnaaagag cagattgega agcccgctgc getegtattg cgagagaaga gtegttactc
                                                                        660
gaagtqccgg gagaggaaaa tgcttgcgag aagaaaqtcg ctggagagaa agccaaqacq
                                                                        720
ttcacgegca tcaagtatge actectcact atgetegaga agtititigga atgegtigee
                                                                        780
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<213> Chlamydia <220> <221> misc_feature 840

897

gacgttttca aattggtgcc gctgcctatt acaatgggta ttcgtgcgat tgtggctgct

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ggatgtacgt tcacttctgc aattattgga ttgtgcactt tctgcgccag agcataa
<210> 264
<211> 298
<212> PRT
<213> Chlamydia
<220>
<221> VARIANT
<222> 202
<223> Xaa = Anv Amino Acid
<400> 264
Met Ala Ser Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu
                                   1.0
Lys Ala Phe Phe Thr Gln Pro Asn Asn Lys Met Ala Arg Val Val Asn
                              25
                                                   30
Lys Thr Lys Gly Val Asp Lys Thr Ile Lys Val Ala Lys Ser Ala Ala
       35
                          40
                                              45
Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser
                       55
                                          60
Ala His Ile Thr Ala Ser Glm Val Ser Lys Gly Leu Gly Asp Ala Arg
                   70
                                       75
Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr
               85
                                  90
Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln
                               105
                                                   110
Lys Thr Gln Glu Gly Asp Glu Gly Leu Thr Ala Asp Leu Cys Val Ser
                          120
His Lys Arg Arg Ala Ala Ala Val Cys Ser Ile Ile Gly Gly Ile
                       135
                                         140
Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
                  150
                                       155
Lys Met Leu Ala Lys Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met
               165
                                   170
Gly Ser Ser Val Ser Tyr Ile Met Ala Ala Asn His Ala Ala Ser Val
           180
                               185
Val Gly Ala Gly Leu Ala Ile Ser Ala Xaa Arg Ala Asp Cys Glu Ala
                           200
                                              205
Arg Cys Ala Arg Ile Ala Arg Glu Glu Ser Leu Leu Glu Val Pro Gly
   210
                                          220
                       215
Glu Glu Asn Ala Cys Glu Lys Lys Val Ala Gly Glu Lys Ala Lys Thr
                   230
                                       235
Phe Thr Arg Ile Lys Tyr Ala Leu Leu Thr Met Leu Glu Lys Phe Leu
               245
                                   250
Glu Cys Val Ala Asp Val Phe Lys Leu Val Pro Leu Pro Ile Thr Met
          260
                              265
                                                  270
Gly Ile Arg Ala Ile Val Ala Ala Gly Cys Thr Phe Thr Ser Ala Ile
       275
                       280
Ile Gly Leu Cys Thr Phe Cys Ala Arg Ala
   290
                       295
<210> 265
<211> 897
<212> DNA
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180

240

300

360

420

480

540

600

660

720

780

840

897

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<222> 604
\langle 223 \rangle n = A.T.C or G
<400> 265
atggetteta tatgeggaeg tttagggtet ggtacaggga atgetetaaa agetttttt
acacagccca acaataaaat ggcaagggta gtaaataaga cgaagggaat ggataagact
attaaggttg ccaagtctgc tgccgaattg accgcaaata ttttggaaca agctggaggc
gegggetett eegcacacat tacagettee caagtgteea aaggattagg ggatgegaga
actgitgtcg ctttagggaa tgcctttaac ggagcgttgc caggaacagt tcaaagtgcg
casagettet teteteacat gasagetget agteagaaaa egeaagaagg ggatgagggg
ctcacagcag atctttgtgt gtctcataag cgcagagcgg ctgcggctgt ctgtagcatc
atoggaggaa ttacctacct cgcgacattc ggagctatcc gtccgattct gtttgtcaac
amantgetgg camaaccgtt tettettee camactamag camatatggg atettetgtt
agotatatta tggcggctaa ccatgcagcg tetgtggtgg gtgetggaet egetateagt
gegnaaagag cagattgega agccegetge getegtattg egagagaaga gtegttacte
gaagtgccgg gagaggaaaa tgcttgcgag aagaaagtcg ctggagagaa agccaagacg
ttcacgegca tcaagtatge actecteact atgetegaga agtttttgga atgegttgee
gacgittica aattggtgcc gctgcctatt acaatgggta ttcgtgcgat tgtggctgct
ggatgtacgt tcacttctgc aattattgga ttgtgcactt tctgcgccag agcataa
<210> 266
<211> 298
<212> PRT
<213> Chlamydia
<220>
<221> VARIANT
<222> 202
<223> Xaa = Any Amino Acid
<400> 266
Met Ala Ser Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu
Lys Ala Phe Phe Thr Gln Pro Asn Asn Lys Met Ala Arg Val Val Asn 20 25 30
Lys Thr Lys Gly Met Asp Lys Thr Ile Lys Val Ala Lys Ser Ala Ala
                           40
Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser
Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Ala Arg
                    70
Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr
Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln
           100
                                105
Lys Thr Gln Glu Gly Asp Glu Gly Leu Thr Ala Asp Leu Cys Val Ser
115 120 125
His Lys Arg Arg Ala Ala Ala Ala Val Cys Ser Ile Ile Gly Gly Ile
                       135
                                        140
Thr Tyr Leu Ala Thr Phe Gly Ala Ile Arg Pro Ile Leu Phe Val Asn
                    150
                                        155
Lys Met Leu Ala Lys Pro Phe Leu Ser Ser Gln Thr Lys Ala Asn Met
                165
                                    170
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Glu Glu Asn Ala Cys Glu Lys Lys Val Ala Gly Glu Lys Ala Lys Thr
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Gly	Ile		260 Ala	Ile	Val	Ala		265 Gly	Cys	Thr	Phe		270 Ser	Ala	Ile	
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Gly Gly Leu Asn Asp Phe His Ala Gly Val Thr Phe Phe Ala Ile Glu Ser Ala Tyr Leu Pro Tyr Thr Val Gln Lys Ser Ser Asp Gly Arg Phe Tyr Phe Val Asp Ile Met Thr Phe Ser Ser Glu Ile Arg Val Gly Asp Glu Leu Leu Glu Val Asp Gly Ala Pro Val Gln Asp Val Leu Ala Thr Leu Tyr Gly Ser Asn His Lys Gly Thr Ala Ala Glu Glu Ser Ala Ala Leu Arg Thr Leu Phe Ser Arg Met Ala Ser Leu Gly His Lys Val Pro Ser Gly Arg Thr Thr Leu Lys Ile Arg Arg Pro Phe Gly Thr Thr Arg 210 215 220 Glu Val Arg Val Lys Trp Arg Tyr Val Pro Glu Gly Val Gly Asp Leu Ala Thr Ile Ala Pro Ser Ile Arg Ala Pro Gln Leu Gln Lys Ser Met Arg Ser Phe Phe Pro Lys Lys Asp Asp Ala Phe His Arg Ser Ser Ser Leu Phe Tyr Ser Pro Met Val Pro His Phe Trp Ala Glu Leu Arg Asn 275 280 285 His Tyr Ala Thr Ser Gly Leu Lys Ser Gly Tyr Asn Ile Gly Ser Thr Asp Gly Phe Leu Pro Val Ile Gly Pro Val Ile Trp Glu Ser Glu Gly Leu Phe Arg Ala Tyr Ile Ser Ser Val Thr Asp Gly Asp Gly Lys Ser 325 330 335 His Lys Val Gly Phe Leu Arg Ile Pro Thr Tyr Ser Trp Gln Asp Met Glu Asp Phe Asp Pro Ser Gly Pro Pro Pro Trp Glu Glu Phe Ala Lys 355 360 365 Ile Ile Gln Val Phe Ser Ser Asn Thr Glu Ala Leu Ile Ile Asp Gln Thr Asn Asn Pro Gly Gly Ser Val Leu Tyr Leu Tyr Ala Leu Leu Ser Met Leu Thr Asp Arg Pro Leu Glu Leu Pro Lys His Arg Met Ile Leu Thr Gln Asp Glu Val Val Asp Ala Leu Asp Trp Leu Thr Leu Leu Glu 425

Asn Val Asp Thr Asn Val Glu Ser Arg Leu Ala Leu Gly Asp Asn Met Glu Gly Tyr Thr Val Asp Leu Gln Val Ala Glu Tyr Leu Lys Ser Phe 455 Gly Arg Gln Val Leu Asn Cys Trp Ser Lys Gly Asp Ile Glu Leu Ser 465 470 475 480 Thr Pro Ile Pro Leu Phe Gly Phe 485 <210> 298 <211> 140 <212> PRT <213> Chlamydia <400> 298 Arg Ile Asp Ile Ser Ser Val Thr Phe Phe Ile Gly Ile Leu Leu Ala Val Asn Ala Leu Thr Tyr Ser His Val Leu Arg Asp Leu Ser Val Ser Met Asp Ala Leu Phe Ser Arg Asn Thr Leu Ala Val Leu Leu Gly Leu Val Ser Ser Val Leu Asp Asn Val Pro Leu Val Ala Ala Thr Ile Gly Met Tyr Asp Leu Pro Met Asn Asp Pro Leu Trp Lys Leu Ile Ala Tyr Thr Ala Gly Thr Gly Gly Ser Ile Leu Ile Ile Gly Ser Ala Ala Gly Val Ala Tyr Met Gly Met Glu Lys Val Ser Phe Gly Trp Tyr Val Lys His Ala Ser Trp Ile Ala Leu Ala Ser Tyr Phe Gly Gly Leu Ala Val Tyr Phe Leu Met Glu Asn Cys Val Asn Leu Phe Val 135 <210> 299

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Asp Ile Pro Ile Val Gly Pro Ser Gly Ser Ala Ala Ser Ala Gly Ser

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Phe	Arg	Ser	Met	11e 85	Glu	Gln	Phe	Asn	Val 90	Asn	Asn	Pro	Ala	Thr 95	Ala
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Arg	Glu	Phe	Val	Asp 325	Gly	Glu	Arg	Ser	Leu 330	Ala	Glu	Ser	Gln	Glu 335	Asn
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Pro Ser Ala Gly Ile Asp Glu Ser Asn Thr Asp Gln Pro Phe Val Leu
Tyr Pro Lys Asp Ile Leu Gly Ser Cys Asn Arg Ile Gly Glu Trp Leu
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Arg Asn Tyr Phe Arg Val Lys Glu Leu Gly Val Ile Ile Thr Asp Ser
His Thr Thr Pro Met Arg Arg Gly Val Leu Gly Ile Gly Leu Cys Trp
Tyr Gly Phe Ser Pro Leu His Asn Tyr Ile Gly Ser Leu Asp Cys Phe
Gly Arg Pro Leu Gln Met Thr Gln Ser Asn Leu Val Asp Ala Leu Ala
Val Ala Ala Val Val Cys Met Gly Glu Gly Asn Glu Gln Thr Pro Leu
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Ala Val Ile Glu Gln Ala Pro Asn Met Val Tyr His Ser Tyr Pro Thr
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Val Thr Thr Asp Val Val Lys Thr Ile Ala Glu Leu Arg Lys Gly Lys Ile Glu Phe Lys Ala Asp Arg Ala Gly Val Cys Asn Val Gly Val Ala Lys Leu Ser Phe Asp Ser Ala Gln Ile Lys Glu Asn Val Glu Ala Leu Cys Ala Ala Leu Val Lys Ala Lys Pro Ala Thr Ala Lys Gly Gln Tyr Leu Val Asn Phe Thr Ile Ser Ser Thr Met Gly Pro Gly Val Thr Val Asp Thr Arg Glu Leu Ile Ala Leu <210> 303 <211> 238 <212> PRT <213> chlamydia <400> 303 Ile Asn Ser Lys Leu Glu Thr Lys Asn Leu Ile Tyr Leu Lys Leu Lys Ile Lys Lys Ser Phe Lys Met Gly Asn Ser Gly Phe Tyr Leu Tyr Asn Thr Gln Asn Cys Val Phe Ala Asp Asn Ile Lys Val Gly Gln Met Thr Glu Pro Leu Lys Asp Gln Gln Ile Ile Leu Gly Thr Thr Ser Thr Pro Val Ala Ala Lys Met Thr Ala Ser Asp Gly Ile Ser Leu Thr Val Ser Asn Asn Pro Ser Thr Asn Ala Ser Ile Thr Ile Gly Leu Asp Ala Glu Lys Ala Tyr Gln Leu Ile Leu Glu Lys Leu Gly Asp Gln Ile Leu Gly Gly Ile Ala Asp Thr Ile Val Asp Ser Thr Val Gln Asp Ile Leu Asp Lys Ile Thr Thr Asp Pro Ser Leu Gly Leu Leu Lys Ala Phe Asn Asn Phe Pro Ile Thr Asn Lys Ile Gln Cys Asn Gly Leu Phe Thr Pro Arg 145 150 155 160

Asn Ile Glu Thr Leu Leu Gly Gly Thr Glu Ile Gly Lys Phe Thr Val 165 175

Thr Pro Lys Ser Ser Gly Ser Met Phe Leu Val Ser Ala Asp Ile Ile Ala Ser Arg Met Glu Gly Gly Val Val Leu Ala Leu Val Arg Glu Gly Asp Ser Lys Pro Tyr Ala Ile Ser Tyr Gly Tyr Ser Ser Gly Val Pro Asn Leu Cys Ser Leu Arg Thr Arg Ile Ile Asn Thr Gly Leu <210> 304 <211> 133 <212> PRT <213> Chlamydia <400> 304 His Met His His His His His His Met Ala Ser Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu Lys Ala Phe Phe Thr Gln Pro Ser Asn Lys Met Ala Arg Val Val Asn Lys Thr Lys Gly Met Asp Lys Thr Val Lys Val Ala Lys Ser Ala Ala Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser Ala His Ile Thr Ala Ser Gln Val Ser Lys Gly Leu Gly Asp Thr Arg Thr Val Val Ala Leu Gly Asn Ala Phe Asn Gly Ala Leu Pro Gly Thr Val Gln Ser Ala Gln Ser Phe Phe Ser His Met Lys Ala Ala Ser Gln Lys Thr Gln Glu Gly Asp Glu Gly 115 120 125 Leu Thr Ala Asp Leu 130 <210> 305 <211> 125 <212> PRT <213> Chlamydia <400> 305 Met Ala Ser Ile Cys Gly Arg Leu Gly Ser Gly Thr Gly Asn Ala Leu Lys Ala Phe Phe Thr Gln Pro Ser Asn Lys Met Ala Arg Val Val Asn Lys Thr Lys Gly Met Asp Lys Thr Val Lys Val Ala Lys Ser Ala Ala Glu Leu Thr Ala Asn Ile Leu Glu Gln Ala Gly Gly Ala Gly Ser Ser 50 60

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1620

1680

1740

1800

1860

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Pro	Gly	Ser	Ser	Leu 245		Leu	Tyr	Thr	Met 250		Ser	Phe	Phe	His 255	
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Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr
Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr
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Ala Met Ala Asp Ala Leu Asn Gly His His Pro Gly Asp Val Ile Ser
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Val Thr Trp Gln Thr Lys Ser Gly Gly Thr Arg Thr Gly Asn Val Thr
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Leu Ala Glu Gly Pro Pro Ala Glu Phe Cys Arg Tyr Pro Ser His Trp
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Arg Pro Leu Asp Pro Val Val Gln Asn Asn Ser Ala Ala Gly Ala Ser
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                    150
Thr Pro Ser Pro Ser Ser Ser Ser Met Pro Gly Ala Val Thr Ile Asn
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Gln Ser Gly Asn Gly Ser Val Ile Phe Thr Ala Glu Ser Leu Thr Pro
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Thr Thr Gly Thr Ile Thr Ala Thr Ser Gly Arg Val Thr Leu Gly Ser
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Gly Ala Ser Leu Ser Ala Val Ala Gly Ala Ala Asn Asn Asn Tyr Thr
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Cys Thr Val Ser Lys Leu Gly Ile Asp Leu Glu Ser Phe Leu Thr Pro 260 265 270
Asn Tyr Lys Thr Ala Ile Leu Gly Ala Asp Gly Thr Val Asn 275 280 285
Ser Gly Ser Thr Leu Asp Leu Val Met Glu Asn Glu Ala Glu Val Tyr
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295

Asp Asn Pro Leu Phe Val Gly Ser Leu Thr Ile Pro Phe Val Thr Leu

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Tyr Gly Glu Tyr Arg Leu Asp Pro Gln Arg Lys Gly Glu Leu Val Pro
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Asn Ser Leu Trp Val Ala Gly Ser Ala Leu Arg Thr Phe Thr Asn Gly
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Leu Lys Glu His Tyr Val Ser Arg Asp Val Gly Phe Val Ala Ser Leu
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His Ala Leu Gly Asp Tyr Ile Leu Asn Tyr Thr Gln Asp Asp Arg Asp
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Gly Phe Leu Ala Arg Tyr Gly Gly Phe Gln Ala Thr Ala Ala Ser His
Tyr Glu Asn Gly Ser Ile Phe Gly Val Ala Phe Gly Gln Leu Tyr Gly
465 470 475 480
Gln Thr Lys Ser Arg Met Tyr Tyr Ser Lys Asp Ala Gly Asn Met Thr
485 490 495
Met Leu Ser Cys Phe Gly Arg Ser Tyr Val Asp Ile Lys Gly Thr Glu 500 505 510
Thr Val Met Tyr Trp Glu Thr Ala Tyr Gly Tyr Ser Val His Arg Met
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His Thr Gln Tyr Phe Asn Asp Lys Thr Gln Lys Phe Asp His Ser Lys
530 540
Cys His Trp His Asn Asn Asn Tyr Tyr Ala Phe Val Gly Ala Glu His
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Asn Phe Leu Glu Tyr Cys Ile Pro Thr Arg Gln Leu Ala Arg Asp Tyr
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Glu Leu Thr Gly Phe Met Arg Phe Glu Met Ala Gly Gly Trp Ser Ser
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Ser Thr Arg Glu Thr Gly Ser Leu Thr Arg Tyr Phe Ala Arg Gly Ser
                600
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Gly His Asn Met Ser Leu Pro Ile Gly Ile Val Ala His Ala Val Ser
610 615 620
His Val Arg Arg Ser Pro Pro Ser Lys Leu Thr Leu Asn Met Gly Tyr 625 630 635
Arg Pro Asp Ile Trp Arg Val Thr Pro His Cys Asn Met Glu Ile Ile
645 650 655
Ala Asn Gly Val Lys Thr Pro Ile Gln Gly Ser Pro Leu Ala Arg His
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Ala Phe Phe Leu Glu Val His Asp Thr Leu Tyr Ile His His Phe Gly
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<212> DNA <213> Chlymadia trachomatis

<400> 330

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<210> 331

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<212> DNA
<213> Chlymadia trachomatis
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accettcata toggectac egectteete geetteggte tigicgacaa caacegeaac
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ggcgcacgag tccaacgcgt ggtcgggagc gctccggcgg caagtctcgg catctccacc
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ggcgacgtga tcaccgcggt cgacggcgct ccgatcaact cggccaccgc gatggcggac
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gegettaaeg ggeateatee eggtgaegte ateteggtga eetggeaaae eaagteggge
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ccatcacact ggoggocget catgaaatgg ctgtcageta ctgcggtgtt tgctgctgtt
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ctcccctcag tttcagggtt ttgcttccca gaacctaaag aattaaattt ctctcgcgta
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gaaacttett cetetaceae ttttaetgaa acaattggag aagetgggge agaatatate
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cocacgaact caaacteete tagetetage ggagaaactg etteegttte tgaggatagt
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totacaatot coctatoagg gattactaaa gogactttot cotgoaacto tgoaqaagtt
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cetgeteetg ttaagaaace tacagaacet aaageteaaa cagcaagega aacgtegggt
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totagtagtt etageggaaa tgatteggtg tettececca gttecagtag agetgaacce
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goagoageta atetteaaag teacttatt tgtgetacag etactectge tgeteaaace
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Phe Leu Gly Leu Gly Val Val Asp Asn Asn Gly Asn Gly Ala Arg Val
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Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr
Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr
Ala Met Ala Asp Ala Leu Asn Gly His His Pro Gly Asp Val Ile Ser
           100
                               105
Val Thr Trp Gln Thr Lys Ser Gly Gly Thr Arg Thr Gly Asn Val Thr
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Arg Pro Leu Met Lys Trp Leu Ser Ala Thr Ala Val Phe Ala Ala Val
145 150 155 160
Leu Pro Ser Val Ser Gly Phe Cys Phe Pro Glu Pro Lys Glu Leu Asn
165 170 175
Phe Ser Arg Val Glu Thr Ser Ser Ser Thr Thr Phe Thr Glu Thr Ile
180 185 190
Gly Glu Ala Gly Ala Glu Tyr Ile Val Ser Gly Asn Ala Ser Phe Thr
195 200 205
Lys Phe Thr Asn Ile Pro Thr Thr Asp Thr Thr Thr Pro Thr Asn Ser 210 215 220
Asn Ser Ser Ser Ser Ser Gly Glu Thr Ala Ser Val Ser Glu Asp Ser 225 230 235
Asp Ser Thr Thr Thr Thr Pro Asp Pro Lys Gly Gly Gly Ala Phe Tyr
245 250 250
Asn Ala His Ser Gly Val Leu Ser Phe Met Thr Arg Ser Gly Thr Glu 260 265 270
Gly Ser Leu Thr Leu Ser Glu Ile Lys Met Thr Gly Glu Gly Gly Ala
275 280 285
Ile Phe Ser Gln Gly Glu Leu Leu Phe Thr Asp Leu Thr Ser Leu Thr 290 300
Ile Gln Asn Asn Leu Ser Gln Leu Ser Gly Gly Ala Ile Phe Gly Gly 305 310 315 320
Ser Thr Ile Ser Leu Ser Gly Ile Thr Lys Ala Thr Phe Ser Cys Asn 325 330 335
Ser Ala Glu Val Pro Ala Pro Val Lys Lys Pro Thr Glu Pro Lys Ala 340 345
Gln Thr Ala Ser Glu Thr Ser Gly Ser Ser Ser Ser Gly Asn Asp 355 360 365
Ser Val Ser Ser Pro Ser Ser Ser Arg Ala Glu Pro Ala Ala Ala Asn
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                                                    380
Leu Gln Ser His Phe Ile Cys Ala Thr Ala Thr Pro Ala Ala Gln Thr
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                                            395
Asp Thr Glu Thr Ser Thr Pro Ser His Lys Pro Gly Ser Gly Gly Ala
405 410 415
Ile Tyr Ala Lys Gly Asp Leu Thr Ile Ala Asp Ser Gln Glu Val Leu 420 425 430
Phe Ser Ile Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu 435 \hspace{1.5cm} 440 \hspace{1.5cm} 445 \hspace{1.5cm}
Lys Asp Val Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn 450 455 460
Gly Ala Glu Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser
465 470 475 480
Ile Gln Ser Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln
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<212> DNA <213> Chlymadia trachomatis

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WO 02/08267 PCT/US01/23121 149

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<213> Chlamydia trachomatis
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<211> 1758
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accepticata toggecetac egecticete ggettgggtg tigtegacaa caacggcaac
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ggcgcacgag tecaacgcgt ggtcgggagc gctccggcgg caagtctcgg catctccacc
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ggcgacgtga tcaccgcggt cgacggcgct ccgatcaact cggccaccgc qatqqcqqac
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gogottaacg ggcatcatcc cggtgacgtc atctcggtga cctggcaaac caagtcgggc
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480
aattotaact acagtaaaca aggtgggggg gototatatg ttgaaggagg tataaactto . 540
caagatottg aagaaattog cattaagtac aataaagotg gaacgttoga aacaaaaaaa
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gotaaaaaag coaagatga cogoataga caactgaa at cototgaga ac otoogctaa 1620
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<211> 585
<212> PRT
<213> Chlamydia trachomatis
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Phe Leu Gly Leu Gly Val Val Asp Asn Asn Gly Asn Gly Ala Arg Val
                      55 60
Gln Arg Val Val Gly Ser Ala Pro Ala Ala Ser Leu Gly Ile Ser Thr
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Gly Asp Val Ile Thr Ala Val Asp Gly Ala Pro Ile Asn Ser Ala Thr

PCT/US01/23121

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Leu	Ala 130	Glu	Gly	Pro	Pro	Ala 135	Glu	Phe	Cys	Arg	Tyr 140	Pro	Ser	His	Trp
Arg 145	Pro	Leu	Gly	Asp	Leu 150	Ser	Ile	Gln	Ser	Ser 155	Lys	Gln	Ser	Leu	Phe 160
Asn	Ser		-	165	_		_	Gly	170			-	Val	Glu 175	Gly
Gly	Ile	Asn	Phe 180	Gln	Asp	Leu	Glu	Glu 185	Ile	Arg	Ile	Lys	Tyr 190	Asn	Lys
Ala	Gly	Thr 195	Phe	Glu	Thr	Lys	Lys 200	Ile	Thr	Leu	Pro	Ser 205	Leu	Lys	Ala
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225			Gly		230					235		-		Ser	240
Gly		-	Ser	245					250					255	_
Gly	_		Tyr 260		-	-		265					270		
Ile		275	Ile				280			-		285	_		
-	Val 290	•	_			295		Glu			300				
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			Leu	325				Gly	330					335	
			Glu 340				_	345			-	_	350	_	
Ala		355	Met				360					365			Thr
	370	-	His	-	_	375					Lys 380				Gln
385	-		Ser	_	390					395					400
_			Val	405		_		Lys	410		Gly			415	
Gly			Thr 420					425					430	Ile	
		435	Asn				440		_	-	_	445		Thr	-
	450		Phe			455	-	Thr			460				
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Ser	Thr		Ser	485	Ser				490	Leu				495	
Ser			500					Ser 505		-			510		
Asn		515	Thr				520	-				525			Ile
ser	530		Thr			535				Ile	540		-	-	
545			Arg		550					555					560
Glu	TTG	GTĀ	Gly	565	1T6	cys	Cys	гля	G1u 570	Ser	ren	Glu	ren	Asp 575	Ala

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<213> Chlamydai trachomatis
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Lys Ser Lys Phe Asp Gln Thr Ala Leu Asp Asn Phe Thr Cys Leu Pro
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65 70 75 80 Pro Ile Glu Val His Glu Arg Glu Ser Ala Lys Gln Gly Arg Glu Val Ser Ala Leu Glu Val Val Leu Thr Val Leu His Ala Gly Gly Lys Phe Asp Lys Asp Ser Tyr Lys Val Ser Gly Gly Leu His Gly Val Gly Val Ser Cys Val Asn Ala Leu Ser Glu Lys Leu Val Ala Thr Val Phe Lys Asp Lys Lys Cys Tyr Gln Met Glu Phe Ser Arg Gly Ile Pro Val Thr Pro Leu Gln Tyr Val Ser Val Ser Asp Arg Gln Gly Thr Glu Ile Val Phe Tyr Pro Asp Pro Lys Ile Phe Ser Thr Cys Thr Phe Asp Arg Ser Ile Leu Met Lys Arg Leu Arg Glu Leu Ala Phe Leu Asn Arg Gly Ile Thr Ile Val Phe Glu Asp Asp Arg Asp Val Ser Phe Asp Lys Val Thr

	210					215					220				
Phe 225	Phe	Tyr	Glu	Gly	Gly 230	Ile	Gln	Ser	Phe	Val 235	Ser	Tyr	Leu	Asn	Gln 240
Asn	Lys	Glu	Ser	Leu 245	Phe	Ser	Glu	Pro	11e 250	Tyr	Ile	Суз	Gly	Thr 255	Arg
Val	Gly	Asp	Asp 260	Gly	Glu	Ile	Glu	Phe 265	Glu	Ala	Ala	Leu	Gln 270	Trp	Asn
Ser	Gly	Tyr 275	Ser	Glu	Leu	Val	Tyr 280	Ser	Tyr	Ala	Asn	Asn 285	Ile	Pro	Thr
Arg	Gln 290	Gly	Gly	Thr	His	Leu 295	Thr	Gly	Phe	Ser	Thr 300	Ala	Leu	Thr	Arg
Val 305	Ile	Asn	Thr	Tyr	11e 310	Lys	Ala	His	Asn	Leu 315	Ala	Lys	Asn	Asn	Lys 320
Leu	Ala	Leu	Thr	Gly 325	Glu	Asp	Ile	Arg	Glu 330	Gly	Leu	Thr	Ala	Val 335	Ile
Ser	Val	Lys	Val 340	Pro	Asn	Pro	Gln	Phe 345	Glu	Gly	Gln	Thr	Lys 350	Gln	Lys
Leu	Gly	Asn 355	Ser	Asp	Val	Ser	Ser 360	Val	Ala	Gln	Gln	Val 365	Val	Gly	Glu
Ala	Leu 370	Thr	Ile	Phe	Phe	Glu 375	Glu	Asn	Pro	Gln	Ile 380	Ala	Arg	Met	Ile
Val 385	Asp	Lys	Val	Phe	Val 390	Ala	Ala	Gln	Ala	Arg 395	Glu	Ala	Ala	Lys	Lys 400
Ala	Arg	Glu	Leu	Thr 405	Leu	Arg	Lys	Ser	Ala 410	Leu	Asp	Ser	Ala	Arg 415	Leu
Pro	Gly	Lys	Leu 420	Ile	Asp	Cys	Leu	Glu 425	Lys	Asp	Pro	Glu	Lys 430	Сув	Glu
Met	Tyr	Ile 435	Val	Glu	Gly	Asp	Ser 440	Ala	Gly	Gly	Ser	Ala 445	Lys	Gln	Gly
Arg	Asp 450	Arg	Arg	Phe	Gln	Ala 455	Ile	Leu	Pro	Ile	Arg 460	Gly	Lys	Ile	Leu
Asn 465	Val	Glu	Lys	Ala	Arg 470	Leu	Gln	Lys	Ile	Phe 475	Gln	Asn	Gln	Glu	Ile 480
Gly	Thr	Ile	Ile	Ala 485	Ala	Leu	Gly	Cys	Gly 490	Ile	Gly	Ala	Asp	Asn 495	Phe
Asn	Leu	Ser	Lys 500	Leu	Arg	Tyr	Arg	Arg 505	Ile	Ile	Ile	Met	Thr 510	Asp	Ala
Asp	Va1	Asp 515	Gly	Ser	His	Ile	Arg 520	Thr	Leu	Leu	Leu	Thr 525	Phe	Phe	Tyr
Arg	His 530	Met	Thr	Ala	Leu	11e 535	Glu	Asn	Glu	Суз	Val 540	Tyr	Ile	Ala	Gln

Pro 545	Pro	Leu	Tyr	Lys	Val 550	Ser	Lys	Lys	Lys	Asp 555	Phe	Arg	Tyr	Ile	Leu 560
Ser	Glu	Lys	Glu	Met 565	Asp	Ser	Tyr	Leu	Leu 570	Met	Leu	Gly	Thr	Asn 575	Glu
Ser	Ser	Ile	Leu 580	Phe	Lys	Ser	Thr	Glu 585	Arg	Glu	Leu	Arg	Gly 590	Glu	Ala
Leu	Glu	Ser 595	Phe	Ile	Asn	Val	11e 600	Leu	Asp	Val	Glu	Ser 605	Phe	Ile	Asn
Thr	Leu 610	Glu	Lys	Lys	Ala	11e 615	Pro	Phe	Ser	Glu	Phe 620	Leu	Glu	Met	Tyr
Lys 625	Glu	Gly	Ile	Gly	Tyr 630	Pro	Leu	Tyr	Tyr	Leu 635	Ala	Pro	Ala	Thr	Gly 640
Met	Gln	Gly	Gly	Arg 645	Tyr	Leu	Tyr	Ser	Asp 650	Glu	G1u	Lys	Glu	Glu 655	Ala
Leu	Ala	Gln	Glu 660	Glu	Thr	His	Lys	Phe 665	Lys	Ile	Ile	Glu	Leu 670	Tyr	Ьуs
Val	Ala	Val 675	Phe	Val	Asp	Ile	Gln 680	Asn	Gl n	Leu	Lys	Glu 685	Tyr	Gly	Leu
Asp	Ile 690	Ser	Ser	Tyr	Leu	Ile 695	Pro	Gln	Lys	Asn	Glu 700	Ile	Val	Ile	Gly
Asn 705	Glu	qsA	Ser	Pro	Ser 710	Cys	Asn	Tyr	Ser	Cys 715	Tyr	Thr	Leu	Glu	Glu 720
Val	Ile	Asn	Tyr	Leu 725	Ьys	Asn	Leu	Gly	Arg 730	Lys	Gly	Ile	Glu	Ile 735	Gln
Arg	Tyr	Lys	Gly 740	Leu	Gly	Glu	Met	Asn 745	Ala	Asp	Gln	Leu	Trp 750	Asp	Thr
Thr	Met	Asn 755	Pro	Glu	Gln	Arg	Thr 760	Leu	Ile	His	Val	Ser 765	Leu	Lys	Asp
Ala	Val 770	Glu	Ala	Asp	His	Ile 775	Phe	Thr	Met	Leu	Met 780	Gly	Glu	Glu	Val
Pro 785	Pro	Arg	Arg	Glu	Phe 790	Ile	Glu	Ser	His	Ala 795	Leu	Ser	Ile	Arg	Ile 800
Asn	Asn	Leu	Asp	11e 805											

<210> 387 <211> 295 <212> PRT <213> Chlamydia pneumoniae

 $<\!400\!>387$ Met Glu Lys Leu Leu Val Thr Asp Ile Asp Gly Thr Ile Thr His Gln 10

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Gln	Ala	Gly 35	Trp	Lys	Leu	Phe	Phe 40	Leu	Thr	Gly	Arg	Tyr 45	Tyr	Lys	Tyr	
Ala	Ala 50	Arg	Leu	Phe	Ser	Asp 55	Phe	Asp	Ala	Pro	Tyr 60	Leu	Leu	Gly	Cys	
Gln 65	Asn	Gly	Ala	Ser	Va1 70	Trp	Ser	Ser	Thr	Ser 75	Ser	Asn	Leu	Leu	Tyr 80	
Ser	Lys	Ser	Leu	Pro 85	Ser	Asp	Leu	Leu	Cys 90	Ile	Leu	Gln	Asp	Cys 95	Met	
Glu	Gly	Ala	Thr 100	Ala	Leu	Phe	Ser	Val 105	Glu	Ser	Gly	Ala	Pro 110	Tyr	G1 y	
Asp	His	Tyr 115	Tyr	Arg	Phe	Ser	Pro 120	Thr	Pro	Ile	Ala	Gln 125	Asp	Leu	His	
Glu	Tyr 130	Val	Asp	Pro	Arg	Tyr 135	Phe	Pro	Aşn	Ala	Lys 140	Glu	Arg	Glu	Ile	
Leu 145	Phe	Glu	Thr	Arg	Ser 150	Leu	Lys	Asp	Asp	Tyr 155	Ala	Phe	Pro	Ser	Phe 160	
Ala	Ala	Ala	Lys	Val 165	Phe	Gly	Leu	Arg	Asp 170	Glu	Val	Ile	Arg	11e 175	Gln	
Lys	G1u	Leu	Glu 180	Arg	Gln	Glu	A1a	Leu 185	Thr	Ser	Val	Ala	Thr 190	Met	Thr	
Leu	Met	Arg 195	Trp	Pro	Phe	Asp	Phe 200	Arg	Tyr	Ala	Ile	Leu 205	Phe	Leu	Thr	
Asp	Lys 210	Ser	Val	Ser	Lys	Gly 215	Lys	Ala	Leu	Asp	Arg 220	Val	Val	Asn	Ile	•
Leu 225	Tyr	Asp	Gly	Lys	Lys 230	Pro	Phe	Val	Met	Ala 235	Ser	Gly	Asp	Asp	Ala 240	
Asn	Asp	Leu	Asp	Leu 245	Ile	Glu	Arg	Gly	Asp 250	Phe	Lys	Ile	Val	Met 255	Ser	
Ser	Ala	Pro	Glu 260	Glu	Met	His	Val	His 265	Ala	Asp	Phe	Leu	Ala 270	Pro	Pro	
Ala	Aap	Lys 275	Asn	Gly	Ile	Leu	Ser 280	Ala	Trp	Glu	Ala	Gly 285	Val	Arg	Tyr	
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<210> 388 <211> 78 <212> PRT <213> Chlamydia pneumoniae

<400> 388

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Ile 1	Thr 210	Leu	Arg	Ala	Ser	11e 215	Asp	Ile	Ile	Asn	Asp 220	Lys	Thr	Leu	Val
Val 1 225	Lys	Gln	Ile	Cys	Pro 230	Gln	Ser	Thr	Thr	Glu 235	Thr	Leu	Ile	Arg	Ser 240
Ile (Glu	Asn	Ala	Ala 245	Lys	Arg	Gly	Thr	11e 250	Lys	Ile	Asp	Thr	Ile 255	Gln
Asp 1	Phe	Ser	Thr 260	Asp	Val	Pro	His	Ile 265	Glu	Ile	Lys	Leu	Pro 270	Lys	Gly
Ser A	Arg	Ala 275	Lys	Glu	Met	Leu	Pro 280	Leu	Leu	Phe	Glu	His 285	Thr	Glu	Cys
Gln Y	Val 290	Ile	Leu	Tyr	Ser	Lys 295	Pro	Thr	Val	Ile	Tyr 300	Glu	Asn	Lys	Pro
Val (Glu	Cys	Ser	Ile	Ser 310	Glu	Ile	Leu	Lys	Leu 315	His	Thr	Thr	Ala	Leu 320
Gln (Gly	Tyr	Leu	Glu 325	Lys	G1u	Leu	Leu	Leu 330	Leu	Gln	Glu	Gln	Leu 335	Thr
Leu A	Asp	His	Tyr 340	His	Lys	Thr	Leu	Glu 345	Tyr	Ile	Phe	Ile	Lys 350	His	Lys
Leu 1	Tyr	Asp 355	Ser	Val	Arg	Glu	Val 360	Leu	Ala	Ile	Asn	Lys 365	Lys	Ile	Ser
Ala A	Asp 370	Asp	Leu	His	Gln	Ala 375	Val	Leu	His	Ala	Leu 380	Glu	Pro	Trp	Leu
His (Glu	Leu	Ala	Thr	Pro 390	Val	Thr	Lys	Gln	Asp 395	Thr	Ser	Gln	Leu	Ala 400
Ser I	Leu	.Thr	Ile	Lys 405	Lys	lle	Leu	Суѕ	Phe 410	Asn	G1u	G1u	Ala	Cys 415	Thr
Lys (Glu	Leu	Leu 420	Ala	Ile	Glu	Lys	Lys 425	Gln	Ala	Ala	Ile	Gln 430	Lys	Asp
Leu (Gly	Arg 435	Ile	Lys	Glu	Val	Thr 440	Val	Lys	Tyr	Leu	Lys 445	Gly	Leu	Leu
Glu A	Arg 450	His	Gly	His	Leu	Gly 455	Glu	Arg	Lys	Thr	Gln 460	Ile	Thr	Asn	Phe
Lys 1 465	Thr	Ala	Lys	Thr	Ser 470	Ile	Leu	Lys	Gln	Gln 475	Thr	Leu	Ile		
<210> <211> <212> <213> <400>	> 25 > PF > Ch	7 T llamy	/dia	pneı	ımoni	Lae									
Met A			Tyr	Ser 5	Pro	Ser	Thr	Ile	Ser 10	Lys	Tyr	Phe	Ile	Tyr 15	Ser

Gly Ala Gly Asn Arg Phe Leu Leu Gly Glu Thr Leu Pro Glu Val Glu Asp Val Arg Phe Leu Cys Gln Glu Thr Arg Val Asp Gly Phe Leu Tyr Leu Lys Pro Ser Ser Cys Ala Asp Ala Gln Leu Ile Ile Phe Asn Ser 50 55 60 Asp Gly Ser Arg Pro Thr Met Cys Gly Asn Gly Leu Arg Cys Ala Ile Ala His Leu Ala Ser Gln Lys Gly Lys Ser Asp Ile Ser Val Ser Thr Asp Ser Gly Leu Tyr Ser Gly Tyr Phe Tyr Ser Trp Asp Arg Val Leu Val Asp Met Thr Leu Ala Asp Trp Arg Ala Ser Val His Arg Leu Glu Ser Arg Pro Asp Pro Leu Pro Lys Glu Val Val Cys Ile His Thr Gly Val Pro His Ala Val Val Ile Leu Pro Glu Ile Ser Thr Leu Asp Leu Ser Ile Leu Gly Pro Phe Leu Arg Tyr His Gln Thr Phe Ser Pro Asp Gly Val Asn Val Asn Phe Val Gln Ile Leu Gly His Cys Gln Leu Arg Val Arg Thr Tyr Glu Arg Gly Val Glu Gly Glu Thr Ala Ala Cys Gly Thr Gly Ala Leu Ala Ser Ala Leu Val Val Ser Asn Ser Tyr Gly Trp Lys Glu Ser Ile Gln Ile His Thr Trp Gly Gly Glu Leu Met Thr Val 225 230 235 240 Ser Gln Asn Arg Gly Arg Val Tyr Leu Gln Gly Ser Val Thr Arg Asp 245 250 255

Leu

<210> 391

<211> 191

<212> PRT

<213> Chlamydia pneumoniae

<400> 391

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Leu Leu Glu Ala Arg Arg Val Phe Phe Ser Glu Pro Val Thr Glu Lys 20 25 30

Ser Ala Ser Asp Ala Ile Lys Lys Leu Trp Tyr Leu Glu Leu Lys Asp Pro Gly Lys Pro Ile Val Phe Val Ile Asn Ser Pro Gly Gly Ser Val 50 $\,$ 60 $\,$ Asp Ala Gly Phe Ala Val Trp Asp Gln Ile Lys Met Leu Thr Ser Pro Val Thr Thr Val Val Thr Gly Leu Ala Ala Ser Met Gly Ser Val Leu Ser Leu Cys Ala Ala Pro Gly Arg Arg Phe Ala Thr Pro His Ser Arg Ile Met Ile His Gln Pro Ser Ile Gly Gly Pro Ile Thr Gly Gln Ala Thr Asp Leu Asp Ile His Ala Arg Glu Ile Leu Lys Thr Lys Ala Arg 130 135 140 Ile Ile Asp Val Tyr Val Glu Ala Thr Asn Gln Pro Arg Asp Ile Ile Glu Lys Ala Ile Asp Arg Asp Met Trp Met Thr Ala Asn Glu Ala Lys Asp Phe Gly Leu Leu Asp Gly Ile Leu Phe Ser Phe Asn Asp Leu 185 <210> 392 <211> 232 <212> PRT <213> Chlamydia pneumoniae <400> 392 Met Thr Lys His Gly Lys Arg Ile Arg Gly Ile Leu Lys Asn Tyr Asp Phe Ser Lys Ser Tyr Ser Leu Arg Glu Ala Ile Asp Ile Leu Lys Gln Cys Pro Pro Val Arg Phe Asp Gln Thr Val Asp Val Ser Ile Lys Leu Gly Ile Asp Pro Lys Lys Ser Asp Gln Gln Ile Arg Gly Ala Val Phe Leu Pro Asn Gly Thr Gly Lys Thr Leu Arg Ile Leu Val Phe Ala Ser Gly Asn Lys Val Lys Glu Ala Val Glu Ala Gly Ala Asp Phe Met Gly Ser Asp Asp Leu Val Glu Lys Ile Lys Ser Gly Trp Leu Glu Phe Asp 100 105 110Val Ala Val Ala Thr Pro Asp Met Met Arg Glu Val Gly Lys Leu Gly

<211> 1723 <212> PRT

<400> 394

<213> Chlamydia pneumoniae

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50 60 Thr Val Glu Gly Asp Leu Arg Arg Arg Val Gln Ser Asp Ile Lys Arg Leu Ile Ala Ile His Ser Tyr Arg Gly Gln Arg His Arg Leu Ser Leu Pro Val Arg Gly Gln Arg Thr Lys Thr Asn Ser Arg Thr Arg Lys Gly Lys Arg Lys Thr Val Ala Gly Lys Lys Lys <210> 394

Met Lys Trp Leu Pro Ala Thr Ala Val Phe Ala Ala Val Leu Pro Ala

				5					10					15	
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Thr	Gly	Ser 35	Gly	Asp	Pro	Thr	Ser 40	Asp	Ala	Ala	Leu	Thr 45	Gly	Phe	Thr
Gln	Ser 50	Ser	Thr	Glu	Thr	Asp 55	Gly	Thr	Thr	Tyr	Thr 60	Ile	Val	Gly	Asp
Ile 65	Thr	Phe	Ser	Thr	Phe 70	Thr	Asn	Ile	Pro	Val 75	Pro	Val	Val	Thr	Pro 80
Asp	Ala	Asn	Asp	Ser 85	Ser	Ser	Asn	Ser	Ser 90	Lys	Gly	Gly	Ser	Ser 95	Ser
Ser	Gly	Ala	Thr 100	Ser	Leu	Ile	Arg	Ser 105	Ser	Asn	Leu	His	Ser 110	Asp	Phe
Asp	Phe	Thr 115	Lys	Asp	Ser	Val	Leu 120	Asp	Leu	Tyr	His	Leu 125	Phe	Phe	Pro
Ser	Ala 130	Ser	Asn	Thr	Leu	Asn 135	Pro	Ala	Leu	Leu	Ser 140	Ser	Ser	Ser	Ser
Gly 145	Gly	Ser	Ser	Ser	Ser 150	Ser	Ser	Ser	Ser	Ser 155	Ser	Gly	Ser	Ala	Ser 160
Ala	Val	Val	Ala	Ala 165	Asp	Pro	Lys	Gly	Gly 170	Ala	Ala	Phe	Tyr	Ser 175	Asn
Glu	Ala	Asn	Gly 180	Thr	Leu	Thr	Phe	Thr 185	Thr	Asp	Ser	Gly	Asn 190	Pro	Gly
Ser	Leu	Thr 195	Leu	Gln	Asn	Leu	Lys 200	Met	Thr	Gly	Asp	Gly 205	Ala	Ala	Ile
Tyr	Ser 210	Lys	Gly	Pro	Leu	Val 215	Phe	Thr	Gly	Leu	Lys 220	Asn	Leu	Thr	Phe
Thr 225	Gly	Asn	Glu	Ser	Gln 230	Lys	Ser	Gly	Gly	Ala 235	Ala	Tyr	Thr	G1 u	Gly 240
Ala	Leu	Thr	Thr	Gln 245	Ala	Ile	Val	Glu	Ala 250	Val	Thr	Phe	Thr	Gly 255	
Thr	Ser	Ala	Gly 260	Gln	Gly	Gly	Ala	11e 265	Tyr	Val	Lys	Glu	Ala 270	Thr	Leu
Phe	Asn	Ala 275	Leu	Asp	Ser	Leu	Lys 280	Phe	Glu	Lys	Asn	Thr 285	Ser	Gly	Gln
Ala	Gly 290	Gly	Gly	Ile	Tyr	Thr 295	Glu	Ser	Thr	Leu	Thr 300	Ile	Ser	Asn	Ile
Thr 305	Lys	Ser	Ile	G l u	Phe 310	Ile	Ser	Asn	Lys	Ala 315	Ser	Val	Pro	Ala	Pro 320
Ala	Pro	Glu	Pro	Thr 325	Ser	Pro	Ala	Pro	Ser 330	Ser	Leu	Ile	Asn	Ser 335	Thr

Thr	Ile	Asp	Thr 340	Ser	Thr	Leu	Gln	Thr 345	Arg	Ala	Ala	Ser	Ala 350	Thr	Pro
Ala	Val	Ala 355	Pro	Val	Ala	Ala	Val 360	Thr	Pro	Thr	Pro	11e 365	Ser	Thr	Gln
Glu	Thr 370	Ala	Gly	Asn	Gly	Gly 375	Ala	Ile	Tyr	Ala	Lys 380	Gln	Gly	Ile	Ser
Ile 385	Ser	Thr	Phe	Lys	Asp 390	Leu	Thr	Phe	Lys	Ser 395	Asn	Ser	Ala	Ser	Val 400
Asp	Ala	Thr	Leu	Thr 405	Val	Asp	Ser	Ser	Thr 410	Ile	Gly	Glu	Ser	GI'y 415	Gly
Ala	Ile	Phe	Ala 420	Ala	Asp	Ser	Ile	Gln 425	Ile	Gln	Gln	Cys	Thr 430	Gly	Thr
Thr	Leu	Phe 435	Ser	Gly	Asn	Thr	Ala 440	Asn	Lys	Ser	Gly	Gly 445	Gly	Ile	Tyr
	450					455					460			Met	
465					470					4/5				Lys	400
				485					490					Asn 495	
			500					505					310	Thr	
		515					520					525		Asn	
	530					535					540			Val	
545					550					555				Ala	560
				565					570					575	
Leu	Thr	Val	Ser 580	Gly	Ile	Thr	Ser	11e 585	Leu	Ser	Phe	Glu	590	Asn	Glu
-		595					600					605		Gln	
	610					615					620			Asp	
625					630					635				ı Leu	640
G1y	Lys	Thr	Leu	Phe 645	Gln	Glu	Asn	Ser	Ser 650	Glu	Lys	His	Gly	655 655	Gly

Leu	Ser	Leu	Ala 660	Ser	Gly	Lys	Ser	Leu 665	Thx	Met	Thr	Ser	Leu 670	Glu	Ser
Phe	Cys	Leu 675	Asn	Ala	Asn	Thx	Ala 680	Lys	Glu	Asn	Gly	Gly 685	Gly	Ala	Asn
Val	Pro 690	Glu	Asn	Ile	Val	Leu 695	Thr	Phe	Thr	Tyr	Thr 700	Pro	Thr	Pro	Asn
Glu 705	Pro	Ala	Pro	Val	Gln 710	Gln	Pro	Val	Tyr	Gly 715	Glu	Ala	Leu	Val	Thr 720
Gly	Asn	Thr	Ala	Thr 725	Lys	Ser	Gly	Gly	Gly 730	Ile	Tyr	Thr	Lys	Asn 735	Ala
Ala	Phe	Ser	Asn 740	Leu	Ser	Ser	Val	Thr 745	Phe	Asp	Gln	Asn	Thr 750	Ser	Ser
Glu	Asn	Gly 755	Gly	Ala	Leu	Leu	Thr 760	Gln	Lys	Ala	Ala	Asp 765	Lys	Thr	Asp
Cys	Ser 770	Phe	Thr	Tyr	Ile	Thr 775	Asn	Val	Asn	Ile	Thr 780	Asn	Asn	Thr	Ala
Thr 785	Gly	Asn	Gly	Gly	Gly 790	Ile	Ala	Gly	Gly	Lys 795	Ala	His	Phe	Asp	Arg 800
Ile	Asp	Asn	Leu	Thr 805	Val	Gln	Ser	Asn	Gln 810	Ala	Lys	Lys	Gly	Gly 815	Gly
Val	Tyr	Leu	Glu 820	Asp	Ala	Leu	Ile	Leu 825	Glu	Lys	Val	Ile	Thr 830	Gly	Ser
Val	Ser	Gln 835	Asn	Thr	Ala	Thr	Glu 840	Ser	Gly	Gly	Gly	Ile 845	Tyr	Ala	Lys
Asp	Ile 850	Gln	Leu	Gln	Ala	Leu 855	Pro	Gly	Ser	Phe	Thr 860	Ile	Thr	Asp	Asn
Lys 865	Val	Glu	Thr	Ser	Leu 870	Thr	Thr	Ser	Thr	Asn 875	Leu	Tyr	Gly	Gly	Gly 880
Ile	Tyr	Ser	Ser	Gly 885	Ala	Val	Thr	Leu	Thr 890	Asn	Ile	Ser	Gly	Thr 895	Phe
Gly	Ile	Thr	Gly 900	Asn	Ser	Val	Ile	Asn 905	Thr	Ala	Thr	Ser	Gln 910	Asp	Ala
Asp	Ile	Gln 915	Gly	Gly	Gly	Ile	Tyr 920	Ala	Thr	Thr	. Ser	Leu 925	Ser	Ile	Asn
Gln	Cys 930	Asn	Thr	Pro	Ile	Leu 935	Phe	Ser	Asn	Asn	Ser 940	Ala	Ala	Thr	Lys
Lys 945	Thr	Ser	Thr	Thr	Lys 950	Gln	Ile	Ala	Gly	Gly 955	Ala	Ile	Phe	Ser	Ala 960
Ala	Val	Thr	Ile	Glu 965	Asn	Asn	Ser	Gln	Pro 970	Ile	Ile	Phe	Leu	975	Asn
Ser	Ala	Lys	Ser	Glu	Ala	Thr	Thr	Ala	Ala	Thr	Ala	Gly	Asn	Lys	Asp

PCT/US01/23121

985 990 980 Ser Cys Gly Gly Ala Ile Ala Ala Asn Ser Val Thr Leu Thr Asn Asn Pro Glu Ile Thr Phe Lys Gly Asn Tyr Ala Glu Thr Gly Gly Ala Ile 1015 Gly Cys Ile Asp Leu Thr Asn Gly Ser Pro Pro Arg Lys Val Ser Ile 1025 Ala Asp Asn Gly Ser Val Leu Phe Gln Asp Asn Ser Ala Leu Asn Arg Gly Gly Ala Ile Tyr Gly Glu Thr Ile Asp Ile Ser Arg Thr Gly Ala 1065 Thr Phe Ile Gly Asn Ser Ser Lys His Asp Gly Ser Ala Ile Cys Cys Ser Thr Ala Leu Thr Leu Ala Pro Asn Ser Gln Leu Ile Phe Glu Asn Asn Lys Val Thr Glu Thr Thr Ala Thr Thr Lys Ala Ser Ile Asn Asn 1115 1110 Leu Gly Ala Ala Ile Tyr Gly Asn Asn Glu Thr Ser Asp Val Thr Ile 1130 Ser Leu Ser Ala Glu Asn Gly Ser Ile Phe Phe Lys Asn Asn Leu Cys 1145 Thr Ala Thr Asn Lys Tyr Cys Ser Ile Ala Gly Asn Val Lys Phe Thr Ala Ile Glu Ala Ser Ala Gly Lys Ala Ile Ser Phe Tyr Asp Ala Val Asn Val Ser Thr Lys Glu Thr Asn Ala Gln Glu Leu Lys Leu Asn Glu Lys Ala Thr Ser Thr Gly Thr Ile Leu Phe Ser Gly Glu Leu His Glu Asn Lys Ser Tyr Ile Pro Gln Lys Val Thr Phe Ala His Gly Asn Leu 1225 Ile Leu Gly Lys Asn Ala Glu Leu Ser Val Val Ser Phe Thr Gln Ser 1240 Pro Gly Thr Thr Ile Thr Met Gly Pro Gly Ser Val Leu Ser Asn His Ser Lys Glu Ala Gly Gly Ile Ala Ile Asn Asn Val Ile Ile Asp Phe Ser Glu Ile Val Pro Thr Lys Asp Asn Ala Thr Val Ala Pro Pro Thr Leu Lys Leu Val Ser Arg Thr Asn Ala Asp Ser Lys Asp Lys Ile Asp 1305

- Ile Thr Gly Thr Val Thr Leu Leu Asp Pro Asn Gly Asn Leu Tyr Gln 1315 1320 1325
- Asn Ser Tyr Leu Gly Glu Asp Arg Asp Ile Thr Leu Phe Asn Ile Asp 1330 1335 1340
- Asn Ser Ala Ser Gly Ala Val Thr Ala Thr Asn Val Thr Leu Gln Gly 1345 1350 1355 136
- Asn Leu Gly Ala Lys Lys Gly Tyr Leu Gly Thr Trp Asn Leu Asp Pro 1365 1370 1375
- Asn Ser Ser Gly Ser Lys Ile Ile Leu Lys Trp Thr Phe Asp Lys Tyr 1380 1380 1390 .
- Leu Arg Trp Pro Tyr Ile Pro Arg Asp Asn His Phe Tyr Ile Asn Ser
- Ile Trp Gly Ala Gln Asn Ser Leu Val Thr Val Lys Gln Gly Ile Leu 1410 1415 1420
- Gly Asn Met Leu Asn Asn Ala Arg Phe Glu Asp Pro Ala Phe Asn Asn 1425 1430 1435
- Phe Trp Ala Ser Ala Ile Gly Ser Phe Leu Arg Lys Glu Val Ser Arg 1445 1445 1450 1450
- Asn Ser Asp Ser Phe Thr Tyr His Gly Arg Gly Tyr Thr Ala Ala Val 1460 1460
- Asp Ala Lys Pro Arg Gln Glu Phe Ile Leu Gly Ala Ala Phe Ser Gln $1475 \hspace{1.5cm} 1480 \hspace{1.5cm} 1485$
- Val Phe Gly His Ala Glu Ser Glu Tyr His Leu Asp Asn Tyr Lys His 1490 1495 1500
- Lys Gly Ser Gly His Ser Thr Gln Ala Ser Leu Tyr Ala Gly Asn Ile 1505 1510 1515 1520
- Phe Tyr Phe Pro Ala Ile Arg Ser Arg Pro Ile Leu Phe Gln Gly Val 1525 Ala Thr Tyr Gly Tyr Met Gln His Asp Thr Thr Tyr Tyr Tyr Pro Ser
- Ala Thr Tyr Gly 1/1 met din in a 1545 1550 1550

 Ile Glu Glu Lys Asn Met Ala Asn Trp Asp Ser Ile Ala Trp Leu Phe
- Asp Leu Arg Phe Ser Val Asp Leu Lys Glu Pro Gln Pro His Ser Thr
- Ala Arg Leu Thr Phe Tyr Thr Glu Ala Glu Tyr Thr Arg Ile Arg Gln
- Glu Lys Phe Thr Glu Leu Asp Tyr Asp Pro Arg Ser Phe Ser Ala Cys 1605 1610 1615
 - Ser Tyr Gly Asn Leu Ala Ile Pro Thr Gly Phe Ser Val Asp Gly Ala 1620 1625 1630

Leu Ala Trp Arg Glu Ile Ile Leu Tyr Asn Lys Val Ser Ala Ala Tyr 1635 1640 1645

Leu Pro Val Ile Leu Arg Asn Asn Pro Lys Ala Thr Tyr Glu Val Leu 1650 1655 1660

Ser Thr Lys Glu Lys Gly Asn Val Val Asn Val Leu Pro Thr Arg Asn 1665 1670 1675 1680

Ala Ala Arg Ala Glu Val Ser Ser Gln Ile Tyr Leu Gly Ser Tyr Trp 1685 1690 1695

Thr Leu Tyr Gly Thr Tyr Thr Ile Asp Ala Ser Met Asn Thr Leu Val 1700 1705 1710

Gln Met Ala Asn Gly Gly Ile Arg Phe Val Phe 1715 1720

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<211> 172.

<213> Chlamydia pneumoniae

<400> 395 Met Lys Trp Leu Pro Ala Thr Ala Val Phe Ala Ala Val Leu Pro Ala 10 Leu Thr Ala Phe Gly Asp Pro Ala Ser Val Glu Ile Ser Thr Ser His 25 Thr Gly Ser Gly Asp Pro Thr Ser Asp Ala Ala Leu Thr Gly Phe Thr 40 Gln Ser Ser Thr Glu Thr Asp Gly Thr Thr Tyr Thr Ile Val Gly Asp 55 60 Ile Thr Phe Ser Thr Phe Thr Asn Ile Pro Val Pro Val Val Thr Pro Asp Ala Asn Asp Ser Ser Ser Asn Ser Ser Lys Gly Gly Ser Ser Ser 90 85 Ser Gly Ala Thr Ser Leu Ile Arg Ser Ser Asn Leu His Ser Asp Phe 100 105 Asp Phe Thr Lys Asp Ser Val Leu Asp Leu Tyr His Leu Phe Phe Pro 125 120 Ser Ala Ser Asn Thr Leu Asn Pro Ala Leu Leu Ser Ser Ser Ser 135 140 150 155 Ala Val Val Ala Ala Asp Pro Lys Gly Gly Ala Ala Phe Tyr Ser Asn 165 170 175 Glu Ala Asn Gly Thr Leu Thr Phe Thr Thr Asp Ser Gly Asn Pro Gly 180 185 Ser Leu Thr Leu Gln Asn Leu Lys Met Thr Gly Asp Gly Ala Ala Ile 195 200 205 Tyr Ser Lys Gly Pro Leu Val Phe Thr Gly Leu Lys Asn Leu Thr Phe 220 215 Thr Gly Asn Glu Ser Gln Lys Ser Gly Gly Ala Ala Tyr Thr Glu Gly 235 230 Ala Leu Thr Thr Gln Ala Ile Val Glu Ala Val Thr Phe Thr Gly Asn 245 250 Thr Ser Ala Gly Gln Gly Gly Ala Ile Tyr Val Lys Glu Ala Thr Leu 270 265 Phe Asn Ala Leu Asp Ser Leu Lys Phe Glu Lys Asn Thr Ser Gly Gln 275 280

Ala Gly Gly Gly Ile Tyr Thr Glu Ser Thr Leu Thr Ile Ser Asn Ile 290 295 300 Thr Lys Ser Ile Glu Phe Ile Ser Asn Lys Ala Ser Val Pro Ala Pro 305 310 315 320 Ala Pro Glu Pro Thr Ser Pro Ala Pro Ser Ser Leu Ile Asn Ser Thr 325 330 Thr Ile Asp Thr Ser Thr Leu Gln Thr Arg Ala Ala Ser Ala Thr Pro 345 Ala Val Ala Pro Val Ala Ala Val Thr Pro Thr Pro Ile Ser Thr Gln 355 360 365 Glu Thr Ala Gly Asn Gly Gly Ala Ile Tyr Ala Lys Gln Gly Ile Ser 370 380 Ile Ser Thr Phe Lys Asp Leu Thr Phe Lys Ser Asn Ser Ala Ser Val 385 390 395 400 Asp Ala Thr Leu Thr Val Asp Ser Ser Thr Ile Gly Glu Ser Gly Gly 405 410 415 Ala Ile Phe Ala Ala Asp Ser Ile Gln Ile Gln Gln Cys Thr Gly Thr 420 425 430 Thr Leu Phe Ser Gly Asn Thr Ala Asn Lys Ser Gly Gly Gly Ile Tyr 435 440 445 435 Ala Val Gly Gln Val Thr Leu Glu Asp Ile Ala Asn Leu Lys Met Thr 450 455 460 Asn Asn Thr Cys Lys Gly Glu Gly Gly Ala Ile Tyr Thr Lys Lys Ala 465 470 475 Leu Thr Ile Asn Asn Gly Ala Ile Leu Thr Thr Phe Ser Gly Asn Thr 485 490 495 Ser Thr Asp Asn Gly Gly Ala Ile Phe Ala Val Gly Gly Ile Thr Leu
500 505 510 Ser Asp Leu Val Glu Val Arg Phe Ser Lys Asn Lys Thr Gly Asn Tyr 515 520 525 Ser Ala Pro Ile Thr Lys Ala Ala Ser Asn Thr Ala Pro Val Val Ser 530 540 Ser Ser Thr Thr Ala Ala Ser Pro Ala Val Pro Ala Ala Ala Ala Ala 545 550 555 560 Pro Val Thr Asn Ala Ala Lys Gly Gly Ala Leu Tyr Ser Thr Glu Gly 565 570 575 Leu Thr Val Ser Gly Ile Thr Ser Ile Leu Ser Phe Glu Asn Asn Glu 580 Cys Gln Asn Gln Gly Gly Gly Ala Tyr Val Thr Lys Thr Phe Gln Cys $\frac{595}{600}$ Ser Asp Ser His Arg Leu Gln Phe Thr Ser Asn Lys Ala Ala Asp Glu 610 615 620 Gly Gly Gly Leu Tyr Cys Gly Asp Asp Val Thr Leu Thr Asn Leu Thr 625 630 635 Gly Lys Thr Leu Phe Gln Glu Asn Ser Ser Glu Lys His Gly Gly Gly 645 655 Leu Ser Leu Ala Ser Gly Lys Ser Leu Thr Met Thr Ser Leu Glu Ser 660 665 670 Phe Cys Leu Asn Ala Asn Thr Ala Lys Glu Asn Gly Gly Gly Ala Asn 675 680 685 Val Pro Glu Asn Ile Val Leu Thr Phe Thr Tyr Thr Pro Thr Pro Asn 690 700 Glu Pro Ala Pro Val Gln Gln Pro Val Tyr Gly Glu Ala Leu Val Thr 705 710 715 720 Gly Asn Thr Ala Thr Lys Ser Gly Gly Gly Ile Tyr Thr Lys Asn Ala 725 730 735 Ala Phe Ser Asn Leu Ser Ser Val Thr Phe Asp Gln Asn Thr Ser Ser 740 745 750 Glu Asn Gly Gly Ala Leu Leu Thr Gln Lys Ala Ala Asp Lys Thr Asp 755 760 765 Cys Ser Fhe Thr Tyr Ile Thr Asn Val Asn Ile Thr Asn Asn Thr Ala

775 Thr Gly Asn Gly Gly Gly Ile Ala Gly Gly Lys Ala His Phe Asp Arg 785 790 795 800 Ile Asp Asn Leu Thr Val Gln Ser Asn Gln Ala Lys Lys Gly Gly Gly 805 810 815 Val Tyr Leu Glu Asp Ala Leu Ile Leu Glu Lys Val Ile Thr Gly Ser 820 825 830 Val Ser Gln Asn Thr Ala Thr Glu Ser Gly Gly Gly Ile Tyr Ala Lys 835 840 845 Asp Ile Gln Leu Gln Ala Leu Pro Gly Ser Phe Thr Ile Thr Asp Asn 850 855 Lys Val Glu Thr Ser Leu Thr Thr Ser Thr Asn Leu Tyr Gly Gly 865 870 875 880 Ile Tyr Ser Ser Gly Ala Val Thr Leu Thr Asn Ile Ser Gly Thr Phe 885 890 895 Gly Ile Thr Gly Asn Ser Val Ile Asn Thr Ala Thr Ser Gln Asp Ala 900 905 910 Asp Ile Gin Gly Gly Gly Ile Tyr Ala Thr Thr Ser Leu Ser Ile Asn 915 920 925 Gln Cys Asn Thr Pro Ile Leu Phe Ser Asn Asn Ser Ala Ala Thr Lys 930 935 940 Lys Thr Ser Thr Thr Lys Gln Ile Ala Gly Gly Ala Ile Phe Ser Ala 950 955 Ala Val Thr Ile Glu Asn Asn Ser Gln Pro Ile Ile Phe Leu Asn Asn 965 970 Ser Ala Lys Ser Glu Ala Thr Thr Ala Ala Thr Ala Gly Asn Lys Asp 985 980 Ser Cys Gly Gly Ala Ile Ala Ala Asn Ser Val Thr Leu Thr Asn Asn 995 1000 1005Pro Glu Ile Thr Phe Lys Gly Asn Tyr Ala Glu Thr Gly Gly Ala Ile 1010 1025 1020 Gly Cys Ile Asp Leu Thr Asn Gly Ser Pro Pro Arg Lys Val Ser Ile 1025 1030 1035 Ala Asp Asn Gly Ser Val Leu Phe Gln Asp Asn Ser Ala Leu Asn Arg 1045 1050 1055 Gly Gly Ala Ile Tyr Gly Glu Thr Ile Asp Ile Ser Arg Thr Gly Ala 1060 1065 1070 Thr Phe Ile Gly Asn Ser Ser Lys His Asp Gly Ser Ala Ile Cys Cys 1075 1080 1085 Ser Thr Ala Leu Thr Leu Ala Pro Asn Ser Gln Leu Ile Phe Glu Asn 1090 1095 1100 Asn Lys Val Thr Glu Thr Thr Ala Thr Thr Lys Ala Ser Ile Asn Asn 1105 1110 1115 1120 Leu Gly Ala Ala Ile Tyr Gly Asn Asn Glu Thr Ser Asp Ile Thr Ile 1125 1130 1135 Ser Leu Ser Ala Glu Asn Gly Ser Ile Phe Phe Lys Asn Asn Leu Cys 1140 1145 1150 Thr Ala Thr Asn Lys Tyr Cys Ser Ile Ala Gly Asn Val Lys Phe Thr 1155 1160 1165 Ala Ile Glu Ala Ser Ala Gly Lys Ala Ile Ser Phe Tyr Asp Ala Val 1170 1175 1180 Asn Val Ser Thr Lys Glu Thr Asn Ala Gln Glu Leu Lys Leu Asn Glu 1185 1190 1195 1200 Lys Ala Thr Ser Thr Gly Thr Ile Leu Phe Ser Gly Glu Leu His Glu 1205 1210 1215 Asn Lys Ser Tyr Ile Pro Gln Lys Val Thr Phe Ala His Gly Asn Leu 1220 1225 1230 Ile Leu Gly Lys Asn Ala Glu Leu Ser Val Val Ser Phe Thr Gln Ser 1235 1240 1245 Pro Gly Thr Thr Ile Thr Met Gly Pro Gly Ser Val Leu Ser Asn His 1250 1255 1260

Ser Lys Glu Ala Gly Gly Ile Ala Ile Asn Asn Val Ile Ile Asp Phe 1265 1270 1275 1280 Ser Glu Ile Val Pro Thr Lys Asp Asn Ala Thr Val Ala Pro Pro Thr 1285 1290 1295 Leu Lys Leu Val Ser Arg Thr Asn Ala Asp Ser Lys Asp Lys Ile Asp 1300 1305 1310 Ile Thr Gly Thr Val Thr Leu Leu Asp Pro Asn Gly Asn Leu Tyr Gln 1315 1320 1325 Asn Ser Tyr Leu Gly Glu Asp Arg Asp Ile Thr Leu Phe Asn Ile Asp 1330 1335 1340 Asn Ser Ala Ser Gly Ala Val Thr Ala Thr Asn Val Thr Leu Gln Gly 1345 1350 1355 1360 Asn Leu Gly Ala Lys Lys Gly Tyr Leu Gly Thr Trp Asn Leu Asp Pro 1365 1370 1375 Asn Ser Ser Gly Ser Lys Ile Ile Leu Lys Trp Thr Phe Asp Lys Tyr 1380 1385 1390 Leu Arg Trp Pro Tyr Ile Pro Arg Asp Asn His Phe Tyr Ile Asn Ser 1395 1400 1405 Ile Trp Gly Ala Gln Asn Ser Leu Val Thr Val Lys Gln Gly Ile Leu
1410 1415 1420 Gly Asn Met Leu Asn Asn Ala Arg Phe Glu Asp Pro Ala Phe Asn Asn 1425 1440 Phe Trp Ala Ser Ala Ile Gly Ser Phe Leu Arg Lys Glu Val Ser Arg 1445 1450 1450 Asn Ser Asp Ser Phe Thr Tyr His Gly Arg Gly Tyr Thr Ala Ala Val 1460 1465 1470 Asp Ala Lys Pro Arg Gln Glu Phe Ile Leu Gly Ala Ala Phe Ser Gln 1475 1480 1485 Val Phe Gly His Ala Glu Ser Glu Tyr His Leu Asp Asn Tyr Lys His 1490 1495 1500 . Lys Gly Ser Gly His Ser Thr Gln Ala Ser Leu Tyr Ala Gly Asn Ile 1505 1510 1520 Phe Tyr Phe Pro Ala Ile Arg Ser Arg Pro Ile Leu Phe Gln Gly Val 1525 1530 1535 Ala Thr Tyr Gly Tyr Met Gln His Asp Thr Thr Thr Tyr Tyr Pro Ser 1540 1545 Ile Glu Glu Lys Asn Met Ala Asn Trp Asp Ser Ile Ala Trp Leu Phe 1555 1560 1565 Asp Leu Arg Phe Ser Val Asp Leu Lys Glu Pro Gln Pro His Ser Thr 1570 1575 1580 Ala Arg Leu Thr Phe Tyr Thr Glu Ala Glu Tyr Thr Arg Ile Arg Gln 1585 1590 1595 Glu Lys Phe Thr Glu Leu Asp Tyr Asp Pro Arg Ser Phe Ser Ala Cys 1605 1610 1615 Ser Tyr Gly Asn Leu Ala Ile Pro Thr Gly Phe Ser Val Asp Gly Ala 1620 1625 1630 Leu Ala Trp Arg Glu Ile Ile Leu Tyr Asn Lys Val Ser Ala Ala Tyr 1635 1640 1645 Leu Pro Val Ile Leu Arg Asn Asn Pro Lys Ala Thr Tyr Glu Val Leu 1650 1660 Ser Thr Lys Glu Lys Gly Asn Val Val Asn Val Leu Pro Thr Arg Asn 1665 1670 1675 168 Ala Ala Arg Ala Glu Val Ser Ser Gln Ile Tyr Leu Gly Ser Tyr Trp 1685 1690 1695 Thr Leu Tyr Gly Thr Tyr Thr Ile Asp Ala Ser Met Asn Thr Leu Val Gln Met Ala Asn Gly Gly Ile Arg Phe Val Phe 1720

<211> 1252 <212> PRT <213> Chlamydia pneumoniae Met Leu Lys Cys Pro Glu Arg Val Ser Val Lys Lys Lys Glu Asp Ile Pro Asp Leu Pro Asn Leu Ile Glu Ile Gln Ile Lys Ser Tyr Lys Gln 20 25 30 Phe Leu Gln Ile Gly Lys Leu Ala Glu Glu Arg Glu Asn Ile Gly Leu Glu Glu Val Phe Arg Glu Ile Phe Pro Ile Lys Ser Tyr Asn Glu Ala 50 60Thr Val Leu Glu Tyr Leu Ser Tyr Asn Leu Gly Val Pro Lys Tyr Ser Pro Glu Glu Cys Ile Arg Arg Gly Ile Thr Tyr Ser Val Thr Leu Lys Val Arg Phe Arg Leu Thr Asp Glu Thr Gly Ile Lys Glu Glu Glu Val Tyr Met Gly Thr Ile Pro Leu Met Thr Asp Lys Gly Thr Phe Ile Ile 125 $$120\$ Asn Gly Ala Glu Arg Val Val Val Ser Gln Val His Arg Ser Pro Gly 130 135 140 Ile Asn Phe Glu Gln Glu Lys His Ser Lys Gly Asn Ile Leu Phe Ser Phe Arg Ile Ile Pro Tyr Arg Gly Ser Trp Leu Glu Ala Ile Phe Asp 165 170 175Ile Asn Asp Leu Ile Tyr Ile His Ile Asp Arg Lys Lys Arg Arg Arg 180 185 190 Lys Ile Leu Ala Ile Thr Phe Ile Arg Ala Leu Gly Tyr Ser Ser Asp 195 200 205 Ala Asp Ile Ile Glu Glu Phe Phe Thr Ile Gly Glu Ser Ser Leu Arg Ser Glu Lys Asp Phe Ala Leu Leu Val Gly Arg Ile Leu Ala Asp Asp 230 235 240 Ile Ile Asp Glu Ala Ser Ser Leu Val Tyr Gly Lys Ala Gly Glu Lys Leu Ser Thr Ala Met Leu Lys Arg Met Leu Asp Ala Gly Ile Ala Ser 260 265 270Val Lys Ile Ala Val Asp Ala Asp Glu Asn His Pro Ile Ile Lys Met Leu Ala Lys Asp Pro Thr Asp Ser Tyr Glu Ala Ala Leu Lys Asp Phe

Tyr 305	Arg	Arg	Leu	Arg	Pro 310	Gly	Glu	Pro	Ala	Thr 315	Leu	Ala	Asn	Ala	Arg 320
Ser	Thr	Ile	Met	Arg 325	Leu	Phe	Phe	Asp	Pro 330	Lys	Arg	Tyr	Asn	Leu 335	Gly
Arg	Val	Gly	Arg 340	Tyr	Lys	Leu	Asn	Arg 345	Lys	Leu	Gly	Phe	Ser 350	Ile	Asp
Asp	Glu	Ala 355	Leu	Ser	Gln	Val	Thr 360	Leu	Arg	Lys	Glu	Asp 365	Val	Ile	Gly
Ala	Leu 370	Lys	Tyr	Leu	Ile	Arg 375	Leu	Lys	Met	Gly	Asp 380	Glu	Lys	Ala	Cys
Val 385	Asp	Asp	Ile	Asp	His 390	Leu	Ala	Asn	Arg	Arg 395	Val	Arg	Ser	Val	Gly 400
Glu	Leu	Ile	Gln	Asn 405	Gln	Cys	Arg	Ser	Gly 410	Leu	Ala	Arg	Met	Glu 415	Lys
Ile	Val	Arg	Glu 420	Arg	Met	Asn	Leu	Phe 425	Asp	Phe	Ser	Ser	Asp 430	Thr	Leu
Thr	Pro	Gly 435	Lys	Val	Val	Ser	Ala 440	Lys	Gly	Leu	Ala	Ser 445	Val	Leu	Lys
Asp	Phe 450	Phe	Gly	Arg	Ser	Gln 455	Leu	Ser	Gln	Phe	Met 460	Asp	Gln	Thr	Asn
Pro 465	Val	Ala	Glu	Leu	Thr 470	His	Lys	Arg	Arg	Leu 475	Ser	Ala	Leu	Gly	Pro 480
Gly	Gly	Leu	Asn	Arg 485	Glu	Arg	Ala	Gly	Phe 490	Glu	Val	Arg	Asp	Val 495	His
Ala	Ser	His	Tyr 500	Gly	Arg	Ile	Cys	Pro 505	Ile	Glu	Thr	Pro	Glu 510	Gly	Pro
Asn	Ile	Gly 515	Leu	Ile	Thr	Ser	Leu 520	Ser	Ser	Phe	Ala	Lys 525	Ile	Asn	Glu
Phe	Gly 530	Phe	Ile	Glu	Thr	Pro 535	Tyr	Arg	Ile	Val	Arg 540	Asp	Gly	Ile	Val
Thr 545	Asp	Glu	Ile	Glu	Tyr 550	Met	Thr	Ala	Asp	Val 555	Glu	Glu	Glu	Cys	Val 560
Ile	Ala	Gln	Ala	Ser 565	Ala	Ser	Leu	Asp	Glu 570	Tyr	Asn	Met	Phe	Thr 575	Glu
Pro	Val	Cys	Trp 580	Val	Arg	Tyr	Ala	Gly 585	Glu	Ala	Phe	Glu	Ala 590	Asp	Thr
Ser	Thr	V al 595	Thr	His	Met	Asp	Val 600	Ser	Pro	Lys	Gln	Leu 605	Val	Ser	Ile
Val	Thr 610	Gly	Leu	Ile	Pro	Phe 615	Leu	Glu	His	Asp	Asp 620	Ala	Asņ	Arg	Ala

Len	Met	Glv	Ser	Asn	Met	Gln	Ara	Gln	Ala	Val	Pro	Len	Leu	Lvs	Thr
625		,			630		9			635				-,,-	640
Glu	Ala	Pro	Val	Val 645	Gly	Thr	Gly	Leu	Glu 650	Cys	Arg	Ala	Ala	Lys 655	Asp
Ser	Gly	Ala	11e 660	Val	Val	Ala	Glu	Glu 665	Asp	Gly	Val	Val	Asp 670	Phe	Val
Asp	Gly	Tyr 675	Lys	Val	Val	Val	Ala 680	Ala	Lys	His	Asn	Pro 685	Thr	Ile	Lys
Arg	Thr 690	Tyr	His	Leu	Lys	Lys 695	Phe	Leu	Arg	Ser	Asn 700	Ser	Gly	Thr	Cys
Ile 705	Asn	Gln	Gln	Pro	Leu 710	Cys	Ala	Val	Gly	Asp 715	Val	Ile	Thr	Lys	Gly 720
Asp	Val	Ile	Ala	Asp 725	Gly	Pro	Ala	Thr	Asp 730	Arg	Gly	Glu	Leu	Ala 735	Leu
Gly	Lys	Asn	Val 740	Leu	Val	Ala	Phe	Met 745	Pro	Trp	Tyr	Gly	Tyr 750	Asn	Phe
Glu	Asp	Ala 755	Ile	Ile	Ile	Ser	Glu 760	Lys	Leu	Ile	Arg	G1u 765	Asp	Ala	Tyr
Thr	Ser 770	Ile	Tyr	Ile	Glu	Glu 775	Phe	Glu	Leu	Thr	Ala 780	Arg	Asp	Thr	Lys
Leu 785	Gly	Lys	Glu	Glu	Ile 790	Thr	Arg	Asp	Ile	Pro 795	Asn	Val	Ser	Asp	Glu 800
Val	Leu	Ala	Asn	Leu 805	Gly	Glu	Asp	Gly	Ile 810	Ile	Arg	Ile	Gly	Ala 815	Glu
Val	Lys	Pro	Gly 820	Asp	Ile	Leu	Val	Gly 825	Lys	Ile	Thr	Pro	Lys 830	Ser	Glu
Thr	Glu	Leu 835	Ala	Pro	Glu	Glu	Arg 840	Leu	Leu	Arg	Ala	Ile 845	Phe	Gly	Glu
Lys	Ala 850	Ala	Asp	Val	Lys	Asp 855	Ala	Ser	Leu	Thr	Val 860	Pro	Pro	Gly	Thr
Glu 865	Gly	Val	Val	Met	Asp 870	Val	Lys	Val	Phe	Ser 875	Arg	Lys	Asp	Arg	Leu 880
Ser	Lys	Ser	Asp	Asp 885	Gl u	Leu	Val	Glu	Glu 890	Ala	Val	His	Leu	Lys 895	Asp
Leu	Gln	Lys	Gly 900	Tyr	Lys	Asn	Gln	Val 905	Ala	Thr	Leu	Lys	Thr 910	G1u	Tyr
Arg	Glu	Lys 915	Leu	Gly	Ala	Leu	Leu 920	Leu	Asn	Glu	Lys	Ala 925	Pro	Ala	Ala
Ile	Ile 930	His	Arg	Arg	Thr	Ala 935	Glu	Ile	Val	Val	His 940	Glu	Gly	Leu	Leu
Phe	Asp	Gln	Glu	Thr	Ile	Glu	Arg	Ile	Glu	Gln	Glu	Asp	Leu	Val	Asp

945		950				955					960
Leu Leu Met	Pro Asn 965	Cys Glu	Met	Tyr	Glu 970	Val	Leu	Lys	Gly	Leu 975	Leu
Ser Asp Tyr	Glu Thr 980	Ala Leu	Gln	Arg 985	Leu	Glu	Ile	Asn	Tyr 990	Lys	Thr
Glu Val Glu 995	His Ile	Arg Glu	Gly 1000		Ala	Asp	Leu	Asp 1005		Gly	Va1
Ile Arg Gln 1010	Val Lys	Val Tyr 101		Ala	Ser	Lys	Arg 1020		Leu	Gln	Val
Gly Asp Lys 1025	Met Ala	Gly Arg 1030	His	Gly	Asn	Lys 1035		Val	Val	Ser	Lys 1040
Ile Val Pro	Glu Ala 104		Pro	Tyr	Leu 1050		Asn	Gly	Glu	Thr 1055	
Gln Met Ile	Leu Asn 1060	Pro Leu	Gly	Val 1065	Pro	Ser	Arg	Met	Asn 1070		Gly
Gln Val Leu 107		His Leu	Gly 1080		Ala	Ala	Lys	Thr 1085		Gly	Ile
Tyr Val Lys 1090	Thr Pro	Val Phe 109		Gly	Phe	Pro	Glu 1100		Arg	Ile	Trp
Asp Met Met 1105	Ile Glu	Gln Gly 1110	Leu	Pro	Glu -	Asp 1115		Lys	Ser	Phe	Leu 1120
Tyr Asp Gly	Lys Thr 112	Gly Glu 5	Arg	Phe	Asp 1130	Asn)	Lys	Val	Val	Ile 1135	
Tyr Ile Tyr	Met Leu 1140	Lys Leu	Ser	His 1145		Ile	Ala	Asp	Lys 1150		His
Ala Arg Ser 115	Ile Gly	Pro Tyr	Ser 1160	Leu)	Val	Thr	Gln	Gln 1165		Leu	Gly
Gly Lys Ala 1170	Gln Met	Gly Gly 117	Gln 5	Arg	Phe	Gly	Glu 1180		Glu	Val	Trp
Ala Leu Glu 1185	Ala Tyr	Gly Val 1190	Ala	His	Met	Leu 1195		Glu	Ile	Leu	Thr 1200
Val Lys Ser				71 50 50	Thr	7	т1 о	Trzw	Gla	Ser	Ile
	Asp Asp 120		GIA	nig	1210		116	Tyr	Ozu	1215	,
Val Lys Gly	120	ō			1210 Gly)				1215 Phe	
Val Lys Gly Val Leu Ile 123	Glu Asn 1220 Lys Glu	beu Leu	Arg	Ser 1225 Leu	Gly	Thr	Pro	Glu	Ser 1230 Arg	1215 Phe	Asn

<211> 224 <212> PRT <213> Chlamydia pneumoniae <400> 397 Met Thr Ser Trp Ile Glu Leu Leu Asp Lys Gln Ile Glu Asp Gln His Met Leu Lys His Glu Phe Tyr Gln Arg Trp Ser Glu Gly Lys Leu Glu 20 25 30 Lys Gln Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys Ala Phe Pro Cys Tyr Leu Ser Ala Leu His Ala Arg Cys Asp Asp Leu
50 55 60 Gln Ile Arg Arg Gln Ile Leu Glu Asn Leu Met Asp Glu Glu Ala Gly Asn Pro Asn His Ile Asp Leu Trp Arg Gln Phe Ala Leu Ser Leu Gly Val Ser Glu Glu Glu Leu Ala Asn His Glu Phe Ser Gln Ala Ala Gln Asp Met Val Ala Thr Phe Arg Arg Leu Cys Asp Met Pro Gln Leu Ala Val Gly Leu Gly Ala Leu Tyr Thr Tyr Glu Ile Gln Ile Pro Gln Val Cys Val Glu Lys Ile Arg Gly Leu Lys Glu Tyr Phe Gly Val Ser Ala 145 150 155 160 Arg Gly Tyr Ala Tyr Phe Thr Val His Gln Glu Ala Asp Ile Lys His Ala Ser Glu Glu Lys Glu Met Leu Gln Thr Leu Val Gly Arg Glu Asn Pro Asp Ala Val Leu Gln Gly Ser Gln Glu Val Leu Asp Thr Leu Trp Asn Phe Leu Ser Ser Phe Ile Asn Ser Thr Glu Pro Cys Ser Cys Lys <210> 398 <211> 556 <212> PRT <213> Chlamydia pneumoniae <400> 398 Met Ser Lys Leu Ile Arg Arg Val Val Thr Val Leu Ala Leu Thr Ser

Met Ala Ser Cys Phe Ala Ser Gly Gly Ile Glu Ala Ala Val Ala Glu
20 25 30

Ser Leu Ile Thr Lys Ile Val Ala Ser Ala Glu Thr Lys Pro Ala Pro

Val Pro Met Thr Ala Lys Lys Val Arg Leu Val Arg Arg Asn Lys Gln Pro Val Glu Gln Lys Ser Arg Gly Ala Phe Cys Asp Lys Glu Phe Tyr
65 70 75 80 Pro Cys Glu Glu Gly Arg Cys Gln Pro Val Glu Ala Gln Gln Glu Ser Cys Tyr Gly Arg Leu Tyr Ser Val Lys Val Asn Asp Asp Cys Asn Val Glu Ile Cys Gln Ser Val Pro Glu Tyr Ala Thr Val Gly Ser Pro Tyr Pro Ile Glu Ile Leu Ala Ile Gly Lys Lys Asp Cys Val Asp Val Val 130 135 140 Ile Thr Gln Gln Leu Pro Cys Glu Ala Glu Phe Val Ser Ser Asp Pro Glu Thr Thr Pro Thr Ser Asp Gly Lys Leu Val Trp Lys Ile Asp Arg 165 170 175 Leu Gly Ala Gly Asp Lys Cys Lys Ile Thr Val Trp Val Lys Pro Leu $180 \hspace{1.5cm} 190 \hspace{1.5cm} 185 \hspace{1.5cm} 190 \hspace{1.5cm}$ Lys Glu Gly Cys Cys Phe Thr Ala Ala Thr Val Cys Ala Cys Pro Glu 195 200 205 Leu Arg Ser Tyr Thr Lys Cys Gly Gln Pro Ala Ile Cys Ile Lys Gln 210 215 220 Glu Gly Pro Asp Cys Ala Cys Leu Arg Cys Pro Val Cys Tyr Lys Ile Glu Val Val Asn Thr Gly Ser Ala Ile Ala Arg Asn Val Thr Val Asp Asn Pro Val Pro Asp Gly Tyr Ser His Ala Ser Gly Gln Arg Val Leu 260 265 270 Ser Phe Asn Leu Gly Asp Met Arg Pro Gly Asp Lys Lys Val Phe Thr Val Glu Phe Cys Pro Gln Arg Arg Gly Gln Ile Thr Asn Val Ala Thr 290 295 300 Val Thr Tyr Cys Gly Gly His Lys Cys Ser Ala Asn Val Thr Thr Val Val Asn Glu Pro Cys Val Gln Val Asn Ile Ser Gly Ala Asp Trp Ser 325 330 335 Tyr Val Cys Lys Pro Val Glu Tyr Ser Ile Ser Val Ser Asn Pro Gly 345 Asp Leu Val Leu His Asp Val Val Ile Gln Asp Thr Leu Pro Ser Gly 360

Val Thr Val Leu Glu Ala Pro Gly Gly Glu Ile Cys Cys Asn Lys Val Val Trp Arg Ile Lys Glu Met Cys Pro Gly Glu Thr Leu Gln Phe Lys Leu Val Val Lys Ala Gln Val Pro Gly Arg Phe Thr Asn Gln Val Ala Val Thr Ser Glu Ser Asn Cys Gly Thr Cys Thr Ser Cys Ala Glu Thr Thr Thr His Trp Lys Gly Leu Ala Ala Thr His Met Cys Val Leu Asp Thr Asn Asp Pro Ile Cys Val Gly Glu Asn Thr Val Tyr Arg Ile Cys 455 Val Thr Asn Arg Gly Ser Ala Glu Asp Thr Asn Val Ser Leu Ile Leu Lys Phe Ser Lys Glu Leu Gln Pro Ile Ala Ser Ser Gly Pro Thr Lys Gly Thr Ile Ser Gly Asn Thr Val Val Phe Asp Ala Leu Pro Lys Leu Gly Ser Lys Glu Ser Val Glu Phe Ser Val Thr Leu Lys Gly Ile Ala Pro Gly Asp Ala Arg Gly Glu Ala Ile Leu Ser Ser Asp Thr Leu Thr Ser Pro Val Ser Asp Thr Glu Asn Thr His Val Tyr <210> 399 <211> 461 <212> PRT <213> Chlamydia pneumoniae <400> 399 Met Thr Gln Glu Phe Asp Cys Val Val Ile Gly Ala Gly Pro Ser Gly Tyr Val Ala Ala Ile Thr Ala Ala Gln Ser Lys Leu Arg Thr Ala Leu Ile Glu Glu Asp Gln Ala Gly Gly Thr Cys Leu Asn Arg Gly Cys Ile Pro Ser Lys Ala Leu Ile Ala Gly Ala Asn Val Val Ser His Ile Lys His Ala Glu Gln Phe Gly Ile His Val Asp Gly Tyr Thr Ile Asp Tyr 65 70 75 80 Pro Ala Met Ala Lys Arg Lys Asn Thr Val Val Gln Gly Ile Arg Gln 85 90 95

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Thr	Gly	Ser 115	Leu	Val	Ser	Ser	Thr 120	Glu	Val	Lys	Val	Ile 125	Gly	Gln	Asp
Thr	Thr 130	Ile	Ile	Lys	Ala	Asn 135	His	Ile	Ile	Leu	Ala 140	Thr	Gly	Ser	Glu
Pro 145	Arg	Pro	Phe	Pro	Gly 150	Val	Pro	Phe	Ser	Ser 155	Arg	Ile	Leu	Ser	Ser 160
Thr	Gly	Ile	Leu	Glu 165	Leu	Glu	Val	Leu	Pro 170	Lys	Lys	Leu	Ala	11e 175	Ile
Gly	Gly	Gly	Val 180	Ile	Gly	Cys	Glu	Phe 185	Ala	Ser	Leu	Phe	His 190	Thr	Leu
Gly	Val	Glu 195	Ile	Thr	Val	Ile	Glu 200	Ala	Leu	Asp	His	Ile 205	Leu	Ala	Val
Asn	Asn 210	Lys	Glu	Val	Ser	Gln 215	Thr	Val	Thr	Asn	Lys 220	Phe	Thr	Lys	Gln
Gly 225	Ile	Arg	Ile	Leu	Thr 230	Lys	Ala	Ser	İle	Ser 235	Ala	Ile	Glu	Glu	Ser 240
Gln	Asn	Gln	Val	Arg 245	Ile	Thr	Val	Asn	Asp 250	Gln	Val	Glu	Glu	Phe 255	Asp
Tyr	Val	Leu	Val 260	Ala	Ile	Gly	Arg	Gln 265	Phe	Asn	Thr	Ala	Ser 270	Ile	Gly
Leu	Asp	Asn 275	Ala	Gly	Val	Ile	Arg 280	Asp	Asp	Arg	Gly	Val 285	Ile	Pro	Val.
Asp	Glu 290	Thr	Met	Arg	Thr	Asn 295	۷al	Pro	Asn	Ile	Tyr 300	Ala	Ile	Gly	Asp
11e 305	Thr	Gly	Lys	Trp	Leu 310	Leu	Ala	His	Val	Ala 315	Ser	His	Gln	Gly	Val 320
Ile	Ala	Ala	Lys	Asn 325	Ile	Ser	Gly	His	His 330	Glu	Val	Met	Asp	Tyr 335	Ser
Ala	Ile	Pro	Ser 340	Val	Ile	Phe	Thr	His 345	Pro	Glu	Ile	Ala	Met 350	Val	Gly
Leu	Ser	Leu 355	Gln	Glu	Ala	Glu	Gln 360	Gln	Asn	Leu	Pro	Ala 365	Lys	Leu	Thr
Lys	Phe 370	Pro	Phe	Lys	Ala	Ile 375	Gly	Lys	Ala	Val	Ala 380	Leu	Gly	Ala	Ser
Asp 385	Gly	Phe	Ala	Ala	Ile 390	Val	Ser	His	Glu	Ile 395	Thr	Gln	Gln	Ile	Leu 400
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Thr Leu Ala Ile Arg Asn Glu Leu Thr Leu Pro Cys Ile Tyr Glu Thr Val His Ala His Pro Thr Leu Ser Glu Val Trp Ala Glu Gly Ala Leu Leu Ala Thr Asn His Pro Leu His Phe Pro Pro Lys Ser 455 <210> 400 <211> 544 <212> PRT <213> Chlamydia pneumoniae <400> 400 Met Ala Ala Lys Asn Ile Lys Tyr Asn Glu Glu Ala Arg Lys Lys Ile His Lys Gly Val Lys Thr Leu Ala Glu Ala Val Lys Val Thr Leu Gly Pro Lys Gly Arg His Val Val Ile Asp Lys Ser Phe Gly Ser Pro Gln Val Thr Lys Asp Gly Val Thr Val Ala Lys Glu Ile Glu Leu Glu Asp Lys His Glu Asn Met Gly Ala Gln Met Val Lys Glu Val Ala Ser Lys Thr Ala Asp Lys Ala Gly Asp Gly Thr Thr Thr Ala Thr Val Leu Ala Glu Ala Ile Tyr Ser Glu Gly Leu Arg Asn Val Thr Ala Gly Ala Asn Pro Met Asp Leu Lys Arg Gly Ile Asp Lys Ala Val Lys Val Val Val 115 120 125 Asp Glu Leu Lys Lys Ile Ser Lys Pro Val Gln His His Lys Glu Ile 130 135 140 Ala Gln Val Ala Thr Ile Ser Ala Asn Asn Asp Ser Glu Ile Gly Asn Leu Ile Ala Glu Ala Met Glu Lys Val Gly Lys Asn Gly Ser Ile Thr 165 170 175 Val Glu Glu Ala Lys Gly Phe Glu Thr Val Leu Asp Val Val Glu Gly Met Asn Phe Asn Arg Gly Tyr Leu Ser Ser Tyr Phe Ser Thr Asn Pro 200 Glu Thr Gln Glu Cys Val Leu Glu Asp Ala Leu Ile Leu Ile Tyr Asp 210 215 . 220 Lys Lys Ile Ser Gly Ile Lys Asp Phe Leu Pro Val Leu Gln Gln Val

Ala	Glu	Ser	Gly	Arg 245	Pro	Leu	Leu	Ile	Ile 250	Ala	Glu	Glu	Ile	Glu 255	Gly
Glu	Ala	Leu	Ala 260	Thr	Leu	Val	Val	Asn 265	Arg	Leu	Arg	Ala	Gly 270	Phe	Arg
Val	Cys	Ala 275	Val	Lys	Ala	Pro	Gly 280	Phe	Gly	Asp	Arg	Arg 285	Lys	Ala	Met
Leu	Glu 290	Asp	Ile	Ala	Ile	Leu 295	Thr	Gly	Gly	Gln	Leu 300	Val	Ser	Glu	Glu
Leu 305	Gly	Met	Lys	Leu	Glu 310	Asn	Thr	Thr	Leu	Ala 315	Met	Leu	Gly	Lys	Ala 320
Lys	Lys	Val	Ile	Val 325	Thr	Lys	Glu	Asp	Thr 330	Thr	Ile	Val	Glu	Gly 335	Leu
Gly	Asn	Lys	Pro 340	Asp	Ile	Gln	Ala	Arg 345	Cys	Asp	Asn	Ile	Lys 350	Lys	Gln
Ile	Glu	Asp 355	Ser	Thr	Ser	Asp	туr 360	Asp	Lys	Glu	Lys	Leu 365	Gln	Glu	Arg
Leu	Ala 370	Lys	Leu	Ser	Gly	Gly 375	Val	Ala	Val	Ile	Arg 380	Val	Gly	Ala	Ala
Thr 385	Glu	Ile	G1u	Met	Lys 390	Glu	Lys	Lys	Asp	Arg 395	Val	Asp	Asp	Ala	Gln 400
His	Ala	Thr	Ile	Ala 405	Ala	Val	Glu	Glu	Gly 410	Ile	Leu	Pro	Gly	Gly 415	Gly
Thr	Ala	Leu	Val 420	Arg	Cys	Ile	Pro	Thr 425	Leu	Glu	Ala	Phe	Leu 430	Pro	Met
Leu	Ala	Asn 435	Glu	Asp	Glu	Ala	11e 440	Gly	Thr	Arg	Ile	Ile 445	Leu	Lys	Ala
Leu	Thr 450	Ala	Pro	Leu	Lys	Gln 455	Ile	Ala	Ser	Asn	Ala 460	Gly	Lys	Glu	Gly
Ala 465	Ile	Ile	Cys	Gln	Gln 470	Val	Leu	Ala	Arg	Ser 475	Ala	Asn	Glu	Gly	Tyr 480
qzA	Ala	Leu	Arg	Asp 485	Ala	Tyr	Thr	Asp	Met 490	Ile	Asp	Ala	Gly	Ile 495	Leu
Asp	Pro	Thr	Lys 500	Val	Thr	Arg	Ser	Ala 505	Leu	Glu	Ser	Ala	Ala 510	Ser	Ile
Ala	Gly	Leu 515	Leu	Leu	Thr	Thr	Glu 520	Ala	Leu	Ile	Ala	Asp 525	Ile	Pro	Glu
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Pro Thr Thr Gln Trp Ile Leu Asp Ile Leu Arg Tyr Trp Val Glu Glu

30	5					310					315					320
Ме	t	His	Val	Asp	Gly 325	Phe	Arg	Phe	Asp	Leu 330	Ala	Ser	Val	Phe	Ser 335	Arg
G1	У	Pro	Ser	Gly 340	Ser	Pro	Leu	Gln	Phe 345	Ala	Pro	Val	Leu	Glu 350	Ala	Ile
Se	er	Phe	Asp 355	Pro	Leu	Leu	Ala	Ser 360	Thr	ьуs	Ile	Ile	Ala 365	Glu	Pro	Trp
As		Ala 370	Gly	Gly	Leu	Tyr	Gln 375	Val	Gly	Tyr	Phe	Pro 380	Thr	Leu	Ser	Pro
Ar 38	g 5	Trp	Ser	G1u	Trp	Asn 390	Gly	Pro	Tyr	Arg	Asp 395	Asn	Val	Lys	Ala	Phe 400
L€	u.	Asn	Gly	Asp	Gln 405	Asn	Leu	Ile	Gly	Thr 410	Phe	Ala	Ser	Arg	I1e 415	Ser
Gl	У	Ser	Gln	Asp 420	Ile	Tyr	Pro	His	Gly 425	Ser	Pro	Thr	Asn	Ser 430	Ile	Asn
Ту	rr '	Val	Ser 435	Cys	His	Asp	Gly	Phe 440	Thr	Leu	Cys	Asp	Thr 445	Val	Thr	Tyr
As	n	His 450	Lys	His	Asn	Glu	Ala 455	Asn	Gly	Gl u	Asp	Asn 460	Arg	Asp	Gly	Thr
As 46		Ala	Asn	Tyr	Ser	Tyr 470	Asn	Phe	Gly	Thr	Glu 475	Gly	Lys	Thr	Glu	Asp 480
Pr	0	Gly	Ile	Leu	Glu 485	Val	Arg	Glu	Arg	Gln 490	Leu	Arg	Asn	Phe	Phe 495	Leu
Th	r	Leu	Met	Val 500	Ser	Gln	Gly	Ile	Pro 505	Met	Ile	Gln	Ser	Gly 510	Asp	Glu
Ту	r.	Ala	His 515	Thr	Ala	Glu	Gly	Asn 520	Asn	Asn	Arg	Trp	Ala 525	Leu	Asp	Ser
As	n.	Ala 530	Asn	Tyr	Phe	Leu	Trp 535	Asp	Gln	Leu	Thr	Ala 540	Lys	Pro	Thr	Leu
Ме 54		His	Phe	Leu	Cys	Asp 550	Leu	Ile	Ala	Phe	Arg 555	Lys	Lys	Tyr	Lys	Thr 560
Le	u :	Phe	Asn	Arg	Gly 565	Phe	Leu	Ser	Asn	Lys 570	Glu	Ile	Ser	Trp	Val 575	Asp
Al	a l	Met	Gly	Asn 580	Pro	Met'	Thr	Trp	Arg 585	Pro	Gly	Asn	Phe	Leu 590	Ala	Phe
Ly	s	Ile	Lys 595	Ser	Pro	Lys	Ala	His 600	Val	Tyr	Val	Ala	Phe 605	His	Val	Gly
Al		Gln 610	Asp	Gln	Leu	Ala	Thr 615	Leu	Pro	Lys	Ala	Ser 620	Ser	Asn	Phe	Leu
Pr 62	5	Tyr	Gln	Ile	Val	Ala 630	Glu	Ser	Gln	Gln	Gly 635	Phe	Val	Pro	Gln	Asn 640

Val Ala Thr Pro Thr Val Ser Leu Gln Pro His Thr Thr Leu Ile Ala $645 \hspace{0.25in} 655 \hspace{0.25in}$

Ile Ser His Ala Lys Glu Val Thr 660

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<211> 328 <212> PRT

<213> Chlamydia pneumoniae

<400> 402

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Gly Val Asp Arg Gly Val Asp Leu Arg Ile Tyr Asp Val Pro Gly Thr 35 40 45

Glu Arg Ala Leu Ser Gly Val Arg Met Glu Leu Asp Asp Gly Ala Tyr 50

Pro Leu Leu His Arg Leu Arg Val Thr Thr Ser Leu Asn Asp Ala Phe 65 70 75 80

Asp Gly Ile Asp Ala Ala Phe Leu Ile Gly Ala Val Pro Arg Gly Pro $85 \hspace{1cm} 90 \hspace{1cm} 95$

Gly Met Glu Arg Gly Asp Leu Leu Lys Gln Asn Gly Gln Ile Phe Ser $100\,$

Leu Gln Gly A1a Ala Leu Asn Thr Ala Ala Lys Arg Asp Ala Lys Ile 115 120 125

Phe Val Val Gly Asn Pro Val Asn Thr Asn Cys Trp Ile Ala Met Lys 130 140 140

His Ala Pro Arg Leu His Arg Lys Asn Phe His Ala Met Leu Arg Leu 145 150 150 160

Asp Gln Asn Arg Met His Ser Met Leu Ala His Arg Ala Glu Val Pro 165 170 175

Leu Glu Glu Val Ser Arg Val Val Ile Trp Gly Asn His Ser Ala Lys 180 180

Gln Val Pro Asp Phe Thr Gln Ala Arg Ile Ser Gly Lys Pro Ala Ala 195 200 205

Glu Val Ile Gly Asp Arg Asp Trp Leu Glu Asn Ile Leu Val His Ser 210 215 220

Val Gln Asn Arg Gly Ser Ala Val Ile Glu Ala Arg Gly Lys Ser Ser 225 230 235 240

Ala Ala Ser Ala Ser Arg Ala Leu Ala Glu Ala Ala Arg Ser Ile Phe

Cys Pro Lys Ser Asp Glu Trp Phe Ser Ser Gly Val Cys Ser Asp His Asn Pro Tyr Gly Ile Pro Glu Asp Leu Ile Phe Gly Phe Pro Cys Arg Met Leu Pro Ser Gly Asp Tyr Glu Ile Ile Pro Gly Leu Pro Trp Glu Pro Phe Ile Arg Asn Lys Ile Gln Ile Ser Leu Asp Glu Ile Ala Gln 305 310 315 Glu Lys Ala Ser Val Ser Ser Leu 325 <210> 403 <211> 217 <212> PRT <213> Chlamydia pneumoniae <400> 403 Met Lys Arg Val Ile Tyr Lys Thr Ile Phe Cys Gly Leu Thr Leu Leu Thr Ser Leu Ser Ser Cys Ser Leu Asp Pro Lys Gly Tyr Asn Leu Glu 20 25 30 Thr Lys Asn Ser Arg Asp Leu Asn Gln Glu Ser Val Ile Leu Lys Glu Asn Arg Glu Thr Pro Ser Leu Val Lys Arg Leu Ser Arg Arg Ser Arg Arg Leu Phe Ala Arg Arg Asp Gln Thr Gln Lys Asp Thr Leu Gln Val Gln Ala Asn Phe Lys Thr Tyr Ala Glu Lys Ile Ser Glu Gln Asp Glu 85 90 95 Arg Asp Leu Ser Phe Val Val Ser Ser Ala Ala Glu Lys Ser Ser Ile Ser Leu Ala Leu Ser Gln Gly Glu Ile Lys Asp Ala Leu Tyr Arg Ile Arg Glu Val His Pro Leu Ala Leu Ile Glu Ala Leu Ala Glu Asn Pro Ala Leu Ile Glu Gly Met Lys Lys Met Gln Gly Arg Asp Trp Ile Trp Asn Leu Phe Leu Thr Gln Leu Ser Glu Val Phe Ser Gln Ala Trp Ser Gln Gly Val Ile Ser Glu Glu Asp Ile Ala Ala Phe Ala Ser Thr Leu 185 Gly Leu Asp Ser Gly Thr Val Ala Ser Ile Val Gln Gly Glu Arg Trp 200

Pro Glu Leu Val Asp Ile Val Ile Thr 210 215 <210> 404 <211> 270 <212> PRT <213> Chlamydia pneumoniae Met Ile Ile Ile Lys Asn Asn Glu Leu Met Ile Arg Arg Phe Phe Lys Thr Leu Phe Pro Pro Gly Pro Gln Tyr Ser Leu Cys Tyr Ala Ser Ile Leu Ile Val Leu Ser Ser Leu Val Cys Val Pro Thr Phe Cys Trp Leu Phe Leu Pro Glu Leu Ser Leu Ser Lys Phe Asn Pro Ser Pro Ile Arg Asn Leu Phe Leu Val Ser Ser Thr Leu Ser Lys Val Pro Pro Thr Ala 65 70 75 80 Ile Ala Glu His Leu Arg Leu Ser Ala Asp Ala Pro Thr Tyr Leu His Glu Phe Ser Ile Lys Glu Ala Glu Ser Ser Leu His Ala Leu Gly Ile Phe Ser Ser Leu Val Ile Glu Lys Ser Pro Asp Asn Lys Gly Ile Thr Ile Phe Tyr Thr Leu Gln Thr Pro Ile Ala Tyr Val Gly Asn Arg Ser Asn Thr Leu Cys Asn Leu Glu Gly Ser Cys Phe Leu Gly Gln Pro Tyr Phe Pro Ser Leu Asn Leu Pro Gln Ile Phe Phe Ser Gln Glu Asp Leu 165 170 175 Lys Met Gln Lys Leu Pro Lys Glu Lys Met Leu Phe Thr Lys Ile Leu Leu Lys Glu Leu Ala Met Glu Ser Pro Lys Ile Ile Asp Leu Ser Leu Ser Asp Ala Tyr Pro Gly Glu Ile Ile Val Thr Leu Ser Ser Gly Ser 210 215 220 Leu Leu Arg Leu Pro Ile Lys Thr Leu Asp Arg Ala Leu Asp Leu Tyr 225 230 235 240 Lys His Met Lys Lys Ser Pro Val Ile Glu Ser Glu Lys Gln Tyr Val

Tyr Asp Leu Arg Phe Pro Asn Phe Leu Leu Leu Lys Ala Leu

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Asp 305	Ser	Pro	Ile	Leu	Gln 310	Glu	Ala	Glu	Gln	Met 315	Val	Ile	Gln	Ala	Glu 320
Lys	Asp	Leu	Lys	Asn 325	Ile	Lys	Pro	Ala	Asp 330	Gly	Ser	Asp	Val	Pro 335	Asn
Pro	Gly	Thr	Thr 340	Val	Gly	Gly	Ser	Lys 345	Gln	Gln	Gly	Ser	Ser 350	Ile	Gly
Ser	Ile	Arg 355	Val	Ser	Met	Leu	Leu 360	Asp	Asp	Ala	Glu	Asn 365	Glu	Thr	Ala
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Glu 385	Asn	Pro	Asp	Ser	Gln 390	Ala	Ala	Gln	Gln	G1u 395	Leu	Ala	Ala	Gln	Ala 400
Arg	Ala	Ala	Lys	Ala 405	Ala	Gly	Asp	Asp	Ser 410	Ala	Ala	Ala	Ala	Leu 415	Ala
Asp	Ala	Gln	Lys 420	Ala	Leu	Glu	Ala	Ala 425	Leu	Gly	Lys	Ala	Gly 430	Gln	Gln
Gln	Gly	Ile 435	Leu	Asn	Ala	Leu	Gly 440	Gln	Ile	Ala	Ser	Ala 445	Ala	Val	Val
Ser	Ala 450	Gly	Val	Pro	Pro	Ala 455	Ala	Ala	Ser	Ser	11e 460	Gly	Ser	Ser	Val
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Gln	Ile	Ser	Ala	Gly 485	Tyr	Asp	Ala	Tyr	Lys 490	Ser	Ile	Asn	Asp	Ala 495	Tyr
Gly	Arg	Ala	Arg 500	Asn	Asp	Ala	Thr	Arg 505	Asp	Val	Ile	Asn	Asn 510	Val	Ser
Thr	Pro	Ala 515	Leu	Thr	Arg	Ser	Val 520	Pro	Arg	Ala	Arg	Thr 525	Glu	Ala	Arg
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Ser 545	Arg	Thr	Leu	Gly	Asp 550	Val	Tyr	Ser	Gln	Val 555	Ser	Ala	Leu	Gln	Ser 560
Val	Met	Gln	Ile	11e 565	Gln	Ser	Asn	Pro	Gln 570	Ala	Asn	Asn	Glu	Glu 575	Ile
Arg	Gln	Lys	Leu 580	Thr	Ser	Ala	Val	Thr 585	Lys	Pro	Pro	Gln	Phe 590	Gly	Tyr
Pro	Туг	Val 595	Gln	Leu	Ser	Asn	Asp 600	Ser	Thr	Gln	Lys	Phe 605	Ile	Ala	Lys

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<213> Chlamydia trachomatis serovar D

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<213> Chlamydia trachomatis serovar D

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<213> Chlamydia trachomatis serovar D

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Ser	Ser 370	Ser	Thr	Ile	His	Asn 375	Ala	Val	Leu	Glu	Ala 380	Leu	Thr	Pro	Phe
Leu 385	Asp	Thr	Leu	Pro	Ala 390	Pro	Asp	Lys	Gln	Ala 395	Thr	Ala	Gln	Leu	Ala 400

Ala Leu Thr Ile Lys Lys Ile Leu Cys Phe Asp Glu Asn Ser Tyr Glu Lys Glu Leu Ala Cys Leu Glu Lys Lys Arg Ser Ser Val Gln Lys Asp 420 425 430 Leu Ser Gln Leu Lys Lys Tyr Thr Val Leu Tyr Ile Lys Lys Leu Leu Glu Thr Tyr Arg Gln Leu Gly His Arg Lys Thr Lys Ile Ala Lys Phe Asp Asp Leu Pro Thr Glu Arg Val Ser Ala His Lys Lys Ala Lys Glu Leu Ala Ala Leu Asp Gln Glu Glu Asn Phe 485 <210> 435 <211> 78 <212> PRT <213> Chlamydia trachomatis serovar D <400> 435 Met Lys Glu Phe Leu Ala Tyr Ile Val Lys Asn Leu Val Asp Lys Pro Glu Glu Val His Leu Lys Glu Val Gln Gly Thr Asn Thr Ile Ile Tyr Glu Leu Thr Val Ala Lys Gly Asp Ile Gly Lys Ile Ile Gly Lys Glu Gly Arg Thr Ile Lys Ala Ile Arg Thr Leu Leu Val Ser Val Ala Ser Arg Asp Asn Val Lys Val Ser Leu Glu Ile Met Glu Glu Arg <210> 436 <211> 647 <212> PRT <213> Chlamydia trachomatis serovar D <400> 436 Met Glu Ser Gly Pro Glu Ser Val Ser Ser Asn Gln Ser Ser Met Asn Pro Ile Ile Asn Gly Gln Ile Ala Ser Asn Ser Glu Thr Lys Glu Ser Thr Lys Glu Ser Glu Ala Ser Pro Ser Ala Ser Ser Ser Val Ser Ser Trp Ser Phe Leu Ser Ser Ala Lys His Ala Leu Ile Ser Leu Arg Asp

Ala Ile Leu Asn Lys Asn Ser Ser Pro Thr Asp Ser Leu Ser Gln Leu

65					70					75					80
Glu	Ala	Ser	Thr	Ser 85	Thr	Ser	Thr	Val	Thr 90	Arg	Val	Ala	Ala	Arg 95	Asp
Tyr	Asn	Glu	Ala 100	Lys	Ser	Asn	Phe	Asp 105	Thr	Ala	Lys	Ser	Gly 110	Leu	Glu
Asn	Ala	Thr 115	Thr	Leu	Ala	Glu	Tyr 120	Glu	Thr	Lys	Met	Ala 125	Asp	Leu	Met
Ala	Ala 130	Leu	Gln	Asp	Met	Glu 135	Arg	Leu	Ala	Lys	Gln 140	Lys	Ala	G1u	Val
Thr 145	Arg	Ile	Lys	Glu	Ala 150	Leu	Gln	Glu	Lys	Gln 155	Glu	Val	Ile	Asp	Lys 160
Leu	Asn	Gln	Leu	Val 165	Lys	Leu	Glu	Lys	Gln 170	Asn	Gln	Thr	Leu	Lys 175	Glu
Thr	Leu	Thr	Thr 180	Thr	Asp	Ser	Ala	Asp 185	Gln	Ile	Pro	Ala	Ile 190	Asn	Ser
Gln	Leu	Glu 195	Ile	Asn	Lys	Asn	Ser 200	Ala	Asp	Gln	Ile	Ile 205	Lys	Asp	Leu
Glu	Gly 210	Gln	Asn	Ile	Ser	Tyr 215	Glu	Ala	Val	Leu	Thr 220	Asn	Ala	Gly	Glu
Val 225	Ile	Lys	Ala	Ser	Ser 230	Glu	Ala	Gly	Ile	Lys 235	Leu	Gly	Gln	Ala	Leu 240
Gln	Ser	Ile	Val	Asp 245	Ala	Gly	Asp	Gln	Ser 250	Gln	Ala	Ala	Val	Leu 255	Gln
Ala	Gln	Gln	Asn 260	Asn	Ser	Pro	Asp	Asn 265	Ile	Ala	Ala	Thr	Lys 270	Lys	Leu
Ile	Asp	Ala 275	Ala	Glu	Thr	Lys	Val 280	Asn	Glu	Leu	Lys	Gln 285	G1u	His	Thr
Gly	Leu 290	Thr	Asp	Ser	Pro	Leu 295	Val	Lys	Lys	Ala	Glu 300	Glu	Gln	Ile	Ser
Gln 305	Ala	Gln	Lys	Asp	Ile 310	Gln	Glu	Ile	Lys	Pro 315	Ser	Gly	Ser	Asp	Ile 320
Pro	Ile	Val	Gly	Pro 325	Ser	Gly	Ser	Ala	Ala 330	Ser	Ala	Gly	Ser	Ala 335	Val
Gly	Ala	Leu	Lys 340	Ser	Ser	Asn	Asn	Ser 345	Gly	Arg	Ile	Ser	Leu 350	Leu	Leu
Asp	Asp	Val 355	Asp	Asn	Glu	Met	Ala 360	Ala	Ile	Ala	Met	Gln 365	Gly	Phe	Arg
Ser	Met 370	Ile	Glu	Gln	Phe	Asn 375	Val	Asn	Asn	Pro	Ala 380	Thr	Ala	Lys	Glu
Leu 385	Gln	Ala	Met	Glu	Ala 390	Gln	Leu	Thr	Ala	Met 395	Ser	Asp	Gln	Leu	Val 400

Gly Ala Asp Gly Glu Leu Pro Ala Glu Ile Gln Ala Ile Lys Asp Ala Leu Ala Gln Ala Leu Lys Gln Pro Ser Thr Asp Gly Leu Ala Thr Ala Met Gly Gln Val Ala Phe Ala Ala Ala Lys Val Gly Gly Gly Ser Ala Gly Thr Ala Gly Thr Val Gln Met Asn Val Lys Gln Leu Tyr Lys Thr Ala Phe Ser Ser Thr Ser Ser Ser Ser Tyr Ala Ala Ala Leu Ser Asp Gly Tyr Ser Ala Tyr Lys Thr Leu Asn Ser Leu Tyr Ser Glu Ser Arg Ser Gly Val Gln Ser Ala Ile Ser Gln Thr Ala Asn Pro Ala Leu Ser Arg Ser Val Ser Arg Ser Gly Ile Glu Ser Gln Gly Arg Ser Ala Asp Ala Ser Gln Arg Ala Ala Glu Thr Ile Val Arg Asp Ser Gln Thr Leu Gly Asp Val Tyr Ser Arg Leu Gln Val Leu Asp Ser Leu Met Ser Thr Ile Val Ser Asn Pro Gln Val Asn Gln Glu Glu Ile Met Gln Lys Leu Thr Ala Ser Ile Ser Lys Ala Pro Gln Phe Gly Tyr Pro Ala Val Gln Asn Ser Ala Asp Ser Leu Gln Lys Phe Ala Ala Gln Leu Glu Arg Glu Phe Val Asp Gly Glu Arg Ser Leu Ala Glu Ser Arg Glu Asn Ala Phe Arg Lys Gln Pro Ala Phe Ile Gln Gln Val Leu Val Asn Ile Ala Ser Leu Phe Ser Gly Tyr Leu Ser

<210> 437

<211> 231

<212> PRT

<213> Chlamydia trachomatis serovar D

<400> 437

Met Met Glu Val Phe Met Asn Phe Leu Asp Gln Leu Asp Leu Ile Ile

Gln Asn Lys His Met Leu Glu His Thr Phe Tyr Val Lys Trp Ser Lys 20 25 30

Gly Glu Leu Thr Lys Glu Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys Ala Phe Pro Lys Tyr Leu Ser Ala Ile His Ser Arg Cys Asp Asp Leu Glu Ala Arg Lys Leu Leu Leu Asp Asn Leu Met Asp Glu Glu Asn Gly Tyr Pro Asn His Ile Asp Leu Trp Lys Gln Phe Val Phe Ala Leu Gly Val Thr Pro Glu Glu Leu Glu Ala His Glu Pro Ser Glu Ala Ala Lys Ala Lys Val Ala Thr Phe Met Arg Trp Cys Thr Gly Asp Ser Leu Ala Ala Gly Val Ala Ala Leu Tyr Ser Tyr Glu Ser Gln Ile Pro Arg Ile Ala Arg Glu Lys Ile Arg Gly Leu Thr Glu Tyr Phe 145 150 155 160 Gly Phe Ser Asn Pro Glu Asp Tyr Ala Tyr Phe Thr Glu His Glu Glu 165 170 175 Ala Asp Val Arg His Ala Arg Glu Glu Lys Ala Leu Ile Glu Met Leu Leu Lys Asp Asp Ala Asp Lys Val Leu Glu Ala Ser Gln Glu Val Thr 195 200 205 Gln Ser Leu Tyr Gly Phe Leu Asp Ser Phe Leu Asp Pro Gly Thr Cys Cys Ser Cys His Gln Ser Tyr 225 230 <210> 438 <211> 533 <212> PRT <213> Chlamydia trachomatis serovar D <400> 438 Met Ser Asn Ser Phe Arg Asp Gln Glu Gln Gly Leu Gln Ala Val Phe Arg Ala Ala Arg Val Ile Ser His Met Phe Ser Gln Thr Ile Gly Pro Tyr Gly Phe Ser Thr Ile Val His Asn Val Gln Asp Thr Arg Thr Thr Gln Asp Ser Gln Ser Met Leu Lys Asp Ile Leu Phe Pro Asp Val Phe Glu Asn Ile Gly Met Lys Leu Ile Arg Asp Thr Ala Leu Arg Thr Arg 65 70 75 80

Met	Arg	Phe	Gly	Asp 85	Gly	Ala	Lys	Thr	Thr 90	Ala	Leu	Leu	Ile	Glu 95	Ala
Leu	Leu	Ala	Glu 100	Gly	Met	Thr	Gly	Ile 105	Gln	Lys	Gly	Leu	Asp 110	Pro	His
Glu	Ile	His 115	Arg	Gly	Met	Leu	Leu 120	Ala	G1u	Lys	Lys	11e 125	Gln	Glu	Val
Phe	Tyr 130	Arg	Glu	Thr	Phe	Pro 135	Leu	Ser	Asp	Leu	Glu 140	His	Thr	Va1	Tyr
Val 145	Ser	Ser	Ile	Ala	Arg 150	Arg	Cys	Asn	Ser	Glu 155	Ile	Ala	Ser	Va1	Leu 160
Ser	Ser	Ala	Val	Gly 165	Tyr	Gly	Gly	Lys	Asn 170	Gly	Tyr	Tyr	Ile	Va1 175	Glu
Glu	His	Glu	Glu 180	His	Glu	Thr	Tyr	Trp 185	His	Ala	Glu	Glu	His 190	Ala	Val
Trp	Asp	Phe 195	Gly	Tyr	Ala	Ser	Pro 200	Tyr	Phe	Ile	Thr	His 205	Ala	Glu	Thr
Gly	Thr 210	Val	Glu	Tyr	Ser	Gln 215		Tyr	Ile	Leu	Val 220	Ser	Glu	Gln	Pro
Leu 225	His	Tyr	Ser	Asn	Pro 230	Ser	Phe	Leu	Thr	Phe 235	Leu	Gln	Ser	Val	Val 240
Gln	Ala	Gly	Lys	Thr 245	Pro	Leu	Val	Ile	Leu 250	Ala	Glu	Ala	Phe	Asp 255	Lys
Glu	Leu	Leu	A1a 260	Met	Leu	Glu	Met	Asn 265	Gln	Ile	Glu	Arg	Val 270	Phe	Pro
Val	Cys	Ala 275	Val	Lys	Val	Ser	Gly 280	Lys	His	Ala	Arg	Glu 285	Ser	Leu	Glu
Asp	11e 290	Ala	Val	Leu	Thr	Gly 295	Ala	Thr	Leu	Leu	Ser 300	Glu	Met	Asp	Phe
Glu 305	Asp	Ser	Glu	Glu	Glu 310	Arg	Ile	Thr	Asn	Arg 315	Leu	Gly	Phe	Val	Ala 320
Gly	Ile	Cys	Val	Ser 325	Ser	Thr	Ser	Leu	Cys 330	Val	Pro	Arg	Glu	Thr 335	Asp
Asn	Lys	Gln	Arg 340	Met	Ala	Glu	His	Cys 345	Ala	Phe	Leu	Gln	Asp 350	Lys	Leu
Ser	Phe	Ser 355	Gln	Glu	Glu	Glu	Ala 360	Ser	Ala	Arg	Leu	Arg 365	Arg	Arg	Leu
Ala	Arg 370	Leu	Ser	Ser	Gly	Glu 375	Val	Cys	Ile	His	Ile 380	Ala	Ala	Asp	Cys
11e 385	Pro	Gln	Glu	Glu	Ile	Gly	Tyr	Ile	Thr	Ser	Ser	Ile	Arg	Ala	Met

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Thr Glu Ser Leu Arg Ser Gly Cys Leu Pro Gly Gly Gly Cys Ala Phe Ile Arg Ala Ala Arg Glu Ile Ser Val Pro Leu Ala Leu Ser Pro Ser Glu Arg Phe Gly Phe Leu Ala Val Leu Ser Ala Ala Glu Lys Pro Phe Arg Ala Ile Val Thr Arg Ser Arg Arg Val Glu Glu Glu Val Phe Ser Glu Val Phe Ser Gln Ala Asp Trp Arg Val Gly Phe Asn Gly Val Ser Gly Phe Val Glu Asp Ile Val Ser Gln Gly Ile Cys Asp Gly Ala Ser Cys Ile Gln Tyr Ala Leu Ser His Ala Val Gly Thr Thr Gly Leu Leu Leu Thr Ser Ala Leu Phe Ile Ala Ser Gln Glu Pro Met Leu Arg Glu Glu Asn Ser Glu Glu 530 <210> 439 <211> 465 <212> PRT <213> Chlamydia trachomatis serovar D <400> 439 Met Asn Glu Ala Phe Asp Cys Val Val Ile Gly Ala Gly Pro Gly Gly
5 10 15 Tyr Val Ala Ala Ile Thr Ala Ala Gln Ala Gly Leu Lys Thr Ala Leu Ile Glu Lys Arg Glu Ala Gly Gly Thr Cys Leu Asn Arg Gly Cys Ile 35 40 45Pro Ser Lys Ala Leu Leu Ala Gly Ala Glu Val Val Thr Gln Ile Arg His Ala Asp Gln Phe Gly Ile His Val Glu Gly Phe Ser Ile Asn Tyr 65 70 75 80Pro Ala Met Val Gln Arg Lys Asp Ser Val Val Arg Ser Ile Arg Asp Gly Leu Asn Gly Leu Ile Arg Ser Asn Lys Ile Thr Val Phe Ser Gly 105 Arg Gly Ser Leu Ile Ser Ser Thr Glu Val Lys Ile Leu Gly Glu Asn

Pro Ser Val Ile Lys Ala His Ser Ile Ile Leu Ala Thr Gly Ser Glu 130 135 140

Lys

Pro 145	Arg	Ala	Phe	Pro	Gly 150	Ile	Pro	Phe	Ser	Ala 155	Glu	Ser	Pro	Arg	11e 160
Leu	Cys	Ser	Thr	Gly 165	Val	Leu	Asn	Leu	Lys 170	Glu	Ile	Pro	Gln	Lys 175	Met
Ala	Ile	Ile	Gly 180	Gly	Gly	Val	Ile	Gly 185	Cys	Glu	Phe	Ala	Ser 190	Leu	Phe
His	Thr	Leu 195	Gly	Ser	Glu	Val	Ser 200	Val	Ile	Glu	Ala	Ser 205	Ser	Gln	Ile
Leu	Ala 210	Leu	Asn	Asn	Pro	Asp 215	Ile	Ser	Lys	Thr	Met 220	Phe	Asp	Lys	Phe
Thr 225	Arg	Gln	Gly	Leu	Arg 230	Phe	Val	Leu	Glu	Ala 235	Ser	Val	Ser	Asn	Ile 240
Glu	Asp	Ile	Gly	Asp 245	Arg	Val	Arg	Leu	Thr 250	Ile	Asn	Gly	Asn	Val 255	Glu
Glu	Tyr	Asp	Tyr 260	Val	Leu	Val	Ser	Ile 265	Gly	Arg	Arg	Leu	Asn 270	Thr	Glu
Asn	Ile	Gly 275	Leu	Asp	Lys	Ala	Gly 280	Val	Ile	Суз	Asp	Glu 285	Arg	Gly	Val
Ile	Pro 290	Thr	Asp	Ala	Thr	Met 295	Arg	Thr	Asn	Val	Pro 300	Asn	Ile	Tyr	Ala
Ile 305	Gly	Asp	Ile	Thr	Gly 310	Lys	Trp	Gln	Leu	Ala 315	His	Val	Ala	Ser	His 320
Gln	Gly	Ile	Ile	Ala 325	Ala	Arg	Asn	Ile	Ala 330	Gly	His	Lys	Glu	Glu 335	Ile
Asp	Tyr	Ser	Ala 340	Val	Pro	Ser	Val	Ile 345	Phe	Thr	Phe	Pro	Glu 350	Val	Ala
Ser	Val	Gly 355	Leu	Ser	Pro	Thr	Ala 360	Ala	Gln	Gln	Gln	Lys 365	Ile	Pro	Val
Lys	Val 370	Thr	Lys	Phe	Pro	Phe 375	Arg	Ala	Ile	Gly	Lys 380	Ala	Val	Ala	Met
Gly 385	Glu	Ala	Asp	Gly	Phe 390	Ala	Ala	Ile	Ile	Ser 395	His	Glu	Thr	Thr	Gln 400
Gln	Ile	Leu	Gly	Ala 405	Tyr	Val	Ile	Gly	Pro 410	His	Ala	Ser	Ser	Leu 415	Ile
Ser	Glu	Ile	Thr 420	Leu	Ala	Val	Arg	Asn 425	Glu	Leu	Thr	Leu	Pro 430	Cys	Ile
Tyr	Glu	Thr 435	Ile	His	Ala	His	Pro 440	Thr	Leu	Ala	Glu	Val 445	Trp	Ala	Glu
Ser	Ala 450	Leu	Leu	Ala	Val	Asp 455	Thr	Pro	Leu	His	Met 460	Pro	Pro	Ala	Lys

<210> 440

<211> 122 <212> PRT

<213> Chlamydia trachomatis serovar D

<400> 440

Met Pro Arg Ile Ile Gly Ile Asp Ile Pro Ala Lys Lys Lys Leu Lys 5 10 15

Ile Ser Leu Thr Tyr Ile Tyr Gly Ile Gly Pro Ala Leu Ser Lys Glu 20 25 30

Ile Ile Ala Arg Leu Gln Leu Asn Pro Glu Ala Arg Ala Ala Glu Leu $35 \ \ 40 \ \ \ 45$

Thr Glu Glu Glu Val Gly Arg Leu Asn Ala Leu Leu Gln Ser Asp Tyr 50 55 60

Val Val Glu Gly Asp Leu Arg Arg Arg Val Gln Ser Asp Ile Lys Arg 65 70 75 80

Leu Ile Thr Ile His Ala Tyr Arg Gly Gln Arg His Arg Leu Ser Leu $85 \hspace{1.5cm} 90 \hspace{1.5cm} 95$

Pro Val Arg Gly Gln Arg Thr Lys Thr Asn Ser Arg Thr Arg Lys Gly $100 \\ 0.05$

Lys Arg Lys Thr Val Ala Gly Lys Lys Lys

<210> 441

<211> 553 <212> PRT

<213> Chlamydia trachomatis serovar D

<400> 441

Met Arg Ile Gly Asp Pro Met Asn Lys Leu Ile Arg Arg Ala Val Thr 5 10 15

Glu Thr Ser Met Ala Glu Ser Leu Ser Thr Asn Val Ile Ser Leu Ala 35 40 45

Asp Thr Lys Ala Lys Asp Asn Thr Ser His Lys Ser Lys Lys Ala Arg 50 60

Lys Asn His Ser Lys Glu Thr Pro Val Asp Arg Lys Glu Val Ala Pro $65 \hspace{1cm} 70 \hspace{1cm} 75 \hspace{1cm} 75 \hspace{1cm} 80 \hspace{1cm}$

Val His Glu Ser Lys Ala Thr Gly Pro Lys Gln Asp Ser Cys Phe Gly $85 \hspace{0.5cm} 90 \hspace{0.5cm} 95$

Arg Met Tyr Thr Val Lys Val Asn Asp Asp Arg Asn Val Glu Ile Thr 100 105 110

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Gln Ala Val Pro Glu Tyr Ala Thr Val Gly Ser Pro Tyr Pro Ile Glu Ile Thr Ala Thr Gly Lys Arg Asp Cys Val Asp Val Ile Ile Thr Gln Gln Leu Pro Cys Glu Ala Glu Phe Val Arg Ser Asp Pro Ala Thr Thr Pro Thr Ala Asp Gly Lys Leu Val Trp Lys Ile Asp Arg Leu Gly Gln 165 170 175 Gly Glu Lys Ser Lys Ile Thr Val Trp Val Lys Pro Leu Lys Glu Gly Cys Cys Phe Thr Ala Ala Thr Val Cys Ala Cys Pro Glu Ile Arg Ser Val Thr Lys Cys Gly Gln Pro Ala Ile Cys Val Lys Gln Glu Gly Pro 210 215 220 Glu Asn Ala Cys Leu Arg Cys Pro Val Val Tyr Lys Ile Asn Ile Val 225 230 235 240 Asn Gln Gly Thr Ala Thr Ala Arg Asn Val Val Val Glu Asn Pro Val Pro Asp Gly Tyr Ala His Ser Ser Gly Gln Arg Val Leu Thr Phe Thr Leu Gly Asp Met Gln Pro Gly Glu His Arg Thr Ile Thr Val Glu Phe Cys Pro Leu Lys Arg Gly Arg Ala Thr Asn Ile Ala Thr Val Ser Tyr 290 295 300 Cys Gly Gly His Lys Asn Thr Ala Ser Val Thr Thr Val Ile Asn Glu Pro Cys Val Gln Val Ser Ile Ala Gly Ala Asp Trp Ser Tyr Val Cys Lys Pro Val Glu Tyr Val Ile Ser Val Ser Asn Pro Gly Asp Leu Val Leu Arg Asp Val Val Val Glu Asp Thr Leu Ser Pro Gly Val Thr Val Leu Glu Ala Ala Gly Ala Gln Ile Ser Cys Asn Lys Val Val Trp Thr Val Lys Glu Leu Asn Pro Gly Glu Ser Leu Gln Tyr Lys Val Leu Val Arg Ala Gln Thr Pro Gly Gln Phe Thr Asn Asn Val Val Lys Ser Cys Ser Asp Cys Gly Thr Cys Thr Ser Cys Ala Glu Ala Thr Thr Tyr 420 425 430Trp Lys Gly Val Ala Ala Thr His Met Cys Val Val Asp Thr Cys Asp

445 435 440 Pro Val Cys Val Gly Glu Asn Thr Val Tyr Arg Ile Cys Val Thr Asn Arg Gly Ser Ala Glu Asp Thr Asn Val Ser Leu Met Leu Lys Phe Ser Lys Glu Leu Gln Pro Val Ser Phe Ser Gly Pro Thr Lys Gly Thr Ile 485 490 495 Thr Gly Asn Thr Val Val Phe Asp Ser Leu Pro Arg Leu Gly Ser Lys Glu Thr Val Glu Phe Ser Val Thr Leu Lys Ala Val Ser Ala Gly Asp Ala Arg Gly Glu Ala Ile Leu Ser Ser Asp Thr Leu Thr Val Pro Val Ser Asp Thr Glu Asn Thr His Ile Tyr <210> 442 <211> 192 <212> PRT <213> Chlamydia trachomatis serovar D Met Pro Glu Gly Glu Met Met His Lys Leu Gln Asp Val Ile Asp Arg Lys Leu-Leu Asp Ser Arg Arg Ile Phe Phe Ser Glu Pro Val Thr Glu Lys Ser Ala Thr Glu Ala Ile Lys Lys Leu Trp Tyr Leu Glu Leu Thr Asn Pro Gly Gln Pro Ile Val Phe Val Ile Asn Ser Pro Gly Gly Ser Val Asp Ala Gly Phe Ala Val Trp Asp Gln Ile Lys Met Ile Ser Ser Pro Leu Thr Thr Val Val Thr Gly Leu Ala Ala Ser Met Gly Ser Val Leu Ser Leu Cys Ala Val Pro Gly Arg Arg Phe Ala Thr Pro His Ala Arg Ile Met Ile His Gln Pro Ser Ile Gly Gly Thr Ile Thr Gly Gln Ala Thr Asp Leu Asp Ile His Ala Arg Glu Ile Leu Lys Thr Lys Ala Arg Ile Ile Asp Val Tyr Val Glu Ala Thr Gly Gln Ser Arg Glu Val Ile Glu Lys Ala Ile Asp Arg Asp Met Trp Met Ser Ala Asn Glu Ala

165

Met Glu Phe Gly Leu Leu Asp Gly Ile Leu Phe Ser Phe Asn Asp Leu 180 185 <210> 443 <211> 275 <212> PRT <213> Chlamydia trachomatis serovar D <400> 443 Met Gly Phe Ser Ser Leu Leu Thr Thr Cys Arg Tyr Leu Leu Tyr Ser Gly Ala Gly Asn Ser Phe Ile Leu Gly Glu Ser Met Pro Ser Leu Glu Asp Val Leu Phe Leu Cys Gln Glu Glu Met Val Asp Gly Phe Leu Cys Val Glu Ser Ser Glu Ile Ala Asp Ala Lys Leu Thr Val Phe Asn Ser 50 55 60 Asp Gly Ser Ile Ala Ser Met Cys Gly Asn Gly Leu Arg Cys Ala Met Ala His Val Ala Gln Cys Phe Gly Leu Glu Asp Val Ser Ile Glu Thr 85 90 95 Glu Arg Gly Val Tyr Gln Gly Lys Phe Phe Ser Met Asn Arg Val Leu Val Asp Met Thr Leu Pro Asp Trp Lys Lys Ala Glu Arg Lys Leu Thr His Val Leu Pro Gly Met Pro Glu Gln Val Phe Phe Ile Asp Thr Gly 135 Val Pro His Val Val Val Phe Val Ser Asp Leu Ser Lys Val Pro Val Gln Glu Trp Gly Ser Phe Leu Arg Tyr His Glu Asp Phe Ala Pro Glu Gly Val Asn Val Asp Phe Val Gln Arg Lys Lys Asp Asp Leu Leu Leu Val Tyr Thr Tyr Glu Arg Gly Cys Glu Arg Glu Thr Leu Ser Cys Gly Thr Gly Met Leu Ala Ser Ala Leu Val Ala Ala Asp Ile Phe Ser Leu Gly Gln Asp Phe Ser Ile Ala Val Cys Ser Arg Ser Arg Asn Leu Ile Lys Ile Phe Ser Glu Lys Gly Lys Val Phe Leu Glu Gly Pro Val Ser Leu Leu Asn Arg Ser Glu Asn Phe Gly Trp Leu Glu Pro Lys Ser Arg WO 02/08267 PCT/US01/23121

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260 265 270 Arg Phe Gly 275 <210> 444 <211> 1770 <212> PRT <213> Chlamydia trachomatis serovar D <400> 444 Met Lys Phe Met Ser Ala Thr Ala Val Phe Ala Ala Ala Leu Ser Ser Val Thr Glu Ala Ser Ser Ile Gln Asp Gln Ile Lys Asn Thr Asp Cys Asn Val Ser Lys Leu Gly Tyr Ser Thr Ser Gln Ala Phe Thr Asp Met Met Leu Ala Asp Asn Thr Glu Tyr Arg Ala Ala Asp Ser Val Ser Phe Tyr Asp Phe Ser Thr Ser Ser Arg Leu Pro Arg Lys His Leu Ser Ser Ser Ser Glu Ala Ser Pro Thr Thr Glu Gly Val Ser Ser Ser Ser Ser Gly Glu Thr Asp Glu Lys Thr Glu Glu Glu Leu Asp Asn Gly Gly Ile Ile Tyr Ala Arg Glu Lys Leu Thr Ile Ser Glu Ser Gln Asp Ser Leu Ser Asn Gln Ser Ile Glu Leu His Asp Asn Ser Ile Phe Phe Gly Glu Gly Glu Val Ile Phe Asp His Arg Val Ala Leu Lys Asn Gly Gly Ala Ile Tyr Gly Glu Lys Glu Val Val Phe Glu Asn Ile Lys Ser Leu Leu Val Glu Val Asn Ile Ala Val Glu Lys Gly Gly Ser Val Tyr Ala Lys Glu Arg Val Ser Leu Glu Asn Val Thr Glu Ala Thr Phe Ser Ser Asn Gly Gly Glu Gln Gly Gly Gly Ile Tyr Ser Glu Gln Asp Met Leu Ile Ser Asp Cys Asn Asn Val His Phe Gln Gly Asn Ala Ala Gly Ala Thr Ala Val Lys Gln Cys Leu Asp Glu Glu Met Ile Val Leu Leu Ala

Glu Cys Val Asp Ser Leu Ser Glu Asp Thr Leu Asp Ser Thr Pro Glu

260 265 Thr Glu Gln Thr Glu Ser Asn Gly Asn Gln Asp Gly Ser Ser Glu Thr Glu Asp Thr Gln Val Ser Glu Ser Pro Glu Ser Thr Pro Ser Pro Asp Asp Val Leu Gly Lys Gly Gly Gly Ile Tyr Thr Glu Lys Ser Leu Thr Ile Thr Gly Ile Thr Gly Thr Ile Asp Phe Val Ser Asn Ile Ala Thr Asp Ser Gly Ala Gly Val Phe Thr Lys Glu Asn Leu Ser Cys Thr Asn 340 350 Thr Asn Ser Leu Gln Phe Leu Lys Asn Ser Ala Gly Gln His Gly Gly Gly Ala Tyr Val Thr Gln Thr Met Ser Val Thr Asn Thr Thr Ser Glu Ser Ile Thr Thr Pro Pro Leu Ile Gly Glu Val Ile Phe Ser Glu Asn Thr Ala Lys Gly His Gly Gly Gly Ile Cys Thr Asn Lys Leu Ser Leu Ser Asn Leu Lys Thr Val Thr Leu Thr Lys Asn Ser Ala Lys Glu Ser Gly Gly Ala Ile Phe Thr Asp Leu Ala Ser Ile Pro Ile Thr Asp Thr Pro Glu Ser Ser Thr Pro Ser Ser Ser Pro Ala Ser Thr Pro Glu Val Val Ala Ser Ala Lys Ile Asn Arg Phe Phe Ala Ser Thr Ala Lys Pro Ala Ala Pro Ser Leu Thr Glu Ala Glu Ser Asp Gln Thr Asp Gln Thr Glu Thr Ser Asp Thr Asn Ser Asp Ile Asp Val Ser Ile Glu Asn 500 505 Ile Leu Asn Val Ala Ile Asn Gln Asn Thr Ser Ala Lys Lys Gly Gly Ala Ile Tyr Gly Lys Lys Ala Lys Leu Ser Arg Ile Asn Asn Leu Glu Leu Ser Gly Asn Ser Ser Gln Asp Val Gly Gly Gly Leu Cys Leu Thr 545 550 555 560 Glu Ser Val Glu Phe Asp Ala Ile Gly Ser Leu Leu Ser His Tyr Asn Ser Ala Ala Lys Glu Gly Gly Ala Ile His Ser Lys Thr Val Thr Leu

Ser Asn Leu Lys Ser Thr Phe Thr Phe Ala Asp Asn Thr Val Lys Ala 600 Ile Val Glu Ser Thr Pro Glu Ala Pro Glu Glu Ile Pro Pro Val Glu Gly Glu Glu Ser Thr Ala Thr Glu Asp Pro Asn Ser Asn Thr Glu Gly Ser Ser Ala Asn Thr Asn Leu Glu Gly Ser Gln Gly Asp Thr Ala Asp Thr Gly Thr Gly Asp Val Asn Asn Glu Ser Gln Asp Thr Ser Asp Thr Gly Asn Ala Glu Ser Glu Glu Gln Leu Gln Asp Ser Thr Gln Ser Asn Glu Glu Asn Thr Leu Pro Asn Ser Asn Ile Asp Gln Ser Asn Glu Asn Thr Asp Glu Ser Ser Asp Ser His Thr Glu Glu Ile Thr Asp Glu Ser Val Ser Ser Ser Glu Ser Gly Ser Ser Thr Pro Gln Asp Gly Gly Ala Ala Ser Ser Gly Ala Pro Ser Gly Asp Gln Ser Ile Ser Ala Asn 740 745 750 Ala Cys Leu Ala Lys Ser Tyr Ala Ala Ser Thr Asp Ser Ser Pro Val Ser Asn Ser Ser Gly Ser Glu Glu Pro Val Thr Ser Ser Ser Asp Ser Asp Val Thr Ala Ser Ser Asp Asn Pro Asp Ser Ser Ser Ser Gly Asp Ser Ala Gly Asp Ser Glu Glu Pro Thr Glu Pro Glu Ala Gly Ser Thr Thr Glu Thr Leu Thr Leu Ile Gly Gly Gly Ala Ile Tyr Gly Glu Thr Val Lys Ile Glu Asn Phe Ser Gly Gln Gly Ile Phe Ser Gly Asn Lys Ala Ile Asp Asn Thr Thr Glu Gly Ser Ser Ser Lys Ser Asp Val Leu Gly Gly Ala Val Tyr Ala Lys Thr Leu Phe Asn Leu Asp Ser Gly Ser Ser Arg Arg Thr Val Thr Phe Ser Gly Asn Thr Val Ser Ser Gln Ser Thr Thr Gly Gln Val Ala Gly Gly Ala Ile Tyr Ser Pro Thr Val Thr

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Ile	Ala	Thr 915	Pro	Val	Val	Phe	Ser 920	Lys	Asn	Ser	Ala	Thr 925	Asn	Asn	Ala
Asn	Asn 930	Thr	Thr	Asp	Thr	G1n 935	Arg	Lys	Asp	Thr	Phe 940	Gly	Gly	Ala	Ile
Gly 945	Ala	Thr	Ser	Ala	Val 950	Ser	Leu	Ser	Gly	Gly 955	Ala	His	Phe	Leu	Glu 960
Asn	Val	Ala	Asp	Leu 965	Gly	Ser	Ala	Ile	Gly 970	Leu	Val	Pro	Gly	Thr 975	Gln
Asn	Thr	Glu	Thr 980	Val	Lys	Leu	Glu	Ser 985	Gly	Ser	Tyr	Tyr	Phe 990	Glu	Lys
Asn	Lys	Ala 995	Leu	Lys	Arg	Ala	Thr 1000		Tyr	Ala	Pro	Val 100		Ser	Ile
Lys	Ala 1010		Thr	Ala	Thr	Phe 101		Gln	Asn	Arg	Ser 1020		Glu	Glu	Gly
Ser 1025		Ile	Tyr	Phe	Thr 1030		Glu	Ala	Ser	Ile 1035		Ser	Leu	Gly	Ser 1040
Val	Leu	Phe	Thr	Gly 1045		Leu	Val	Thr	Leu 1050		Leu	Ser	Thr	Thr 105	
Glu	Gly	Thr	Pro 1060		Thr	Thr	Ser	Gly 1065		Val	Thr	Lys	Tyr 1070		Ala
Ala	Ile	Phe 1075	Gly	Gln	Ile	Ala	Ser 1080		Asn	Gly	Ser	Gln 108		Asp	Asn
Leu	Pro 1090		Lys	Leu	Ile	Ala 1095		Gly	Gly	Asn	Ile 1100		Phe	Arg	Asn
Asn 1105		Tyr	Arg	Pro	Thr 1110		Ser	Asp	Thr	Gly 1115		Ser	Thr	Phe	Cys 1120
Ser	Ile	Ala	Gly	Asp 1125		Lys	Leu	Thr	Met 1130		Ala	Ala	Lys	Gly 1135	
Thr	Ile	Ser	Phe 1140		Asp	Ala	Ile	Arg 1145		Ser	Thr	Lys	Lys 1150		Gly
Thr	Gln	Ala 1155	Thr	Ala	Tyr	Asp	Thr 1160		Asp	Ile	Asn	Lys 116		Glu	Asp
Ser	Glu 1170		Val	Asn	Ser	Ala 1175		Thr	Gly	Thr	Ile 1180		Phe	Ser	Ser
Glu 1185		His	Glu	Asn	Lys 1190		Tyr	Ile	Pro	Gln 1195		Val	Va1	Leu	His 1200
Ser	Gly	Ser	Leu	Val 1205		Lys	Pro	Asn	Thr 1210		Leu	His	Val	Ile 1215	
Phe	Glu	Gln	Lys 1220		Gly	Ser	Ser	Leu 1225		Met	Thr	Pro	Gly 1230		Val
Leu	Ser	Asn	Gln	Thr	Va1	Ala	Asp	Gly	Ala	Leu	Val	Ile	Asn	Asn	Met

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1235 1240 1245 Thr Ile Asp Leu Ser Ser Val Glu Lys Asn Gly Ile Ala Glu Gly Asn 1255 1260 Ile Phe Thr Pro Pro Glu Leu Arg Ile Ile Asp Thr Thr Thr Gly Gly Ser Gly Gly Thr Pro Ser Thr Asp Ser Glu Ser Asn Gln Asn Ser Asp 1285 1290 Asp Thr Glu Glu Gln Asn Asn Asp Ala Ser Asn Gln Gly Glu Ser 1305 Ala Asn Gly Ser Ser Ser Pro Ala Val Ala Ala Ala His Thr Ser Arg Thr Arc Asn Phe Ala Ala Ala Ala Thr Ala Thr Pro Thr Thr Thr Pro 1335 1340 Thr Ala Thr Thr Thr Ser Asn Gln Val Ile Leu Gly Gly Glu Ile Lys Leu Ile Asp Pro Asn Gly Thr Phe Phe Gln Asn Pro Ala Leu Arg 1365 Ser Asp Gln Gln Ile Ser Leu Leu Val Leu Pro Thr Asp Ser Ser Lys 1385 Met Gln Ala Gln Lys Ile Val Leu Thr Gly Asp Ile Ala Pro Gln Lys Gly Tyr Thr Gly Thr Leu Thr Leu Asp Pro Asp Gln Leu Gln Asn Gly 1415 Thr Ile Ser Val Leu Trp Lys Phe Asp Ser Tyr Arg Gln Trp Ala Tyr Val Pro Arg Asp Asn His Phe Tyr Ala Asn Ser Ile Leu Gly Ser Gln Met Leu Met Val Thr Val Lys Gln Gly Leu Leu Asn Asp Lys Met Asn 1465 Leu Ala Arq Phe Glu Glu Val Ser Tyr Asn Asn Leu Trp Ile Ser Gly Leu Gly Thr Met Leu Ser Gln Val Gly Thr Pro Thr Ser Glu Glu Phe 1495 Thr Tyr Tyr Ser Arg Gly Ala Ser Val Ala Leu Asp Ala Lys Pro Ala His Asp Val Ile Val Gly Ala Ala Phe Ser Lys Met Ile Gly Lys Thr Lys Ser Leu Lys Arg Glu Asn Asn Tyr Thr His Lys Gly Ser Glu Tyr 1545 Ser Tyr Gln Ala Ser Val Tyr Gly Gly Lys Pro Phe His Phe Val Ile 1555 1560 1565

- Asn Lys Lys Thr Glu Lys Ser Leu Pro Leu Leu Leu Gln Gly Val Ile $1570 \hspace{1.5cm} 1575 \hspace{1.5cm} 1580$
- Ser Tyr Gly Tyr Ile Lys His Asp Thr Val Thr His Tyr Pro Thr Ile 1585 1590 1595 1600
- Arg Glu Arg Asn Lys Gly Glu Trp Glu Asp Leu Gly Trp Leu Thr Ala 1605 1610 1615
- Leu Arg Val Ser Ser Val Leu Arg Thr Pro Ala Gln Gly Asp Thr Lys $1620 \hspace{1.5cm} 1625 \hspace{1.5cm} 1630$
- Arg Ile Thr Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys 1635 1640 1645
- Gln Phe Thr Glu Thr Glu Tyr Asp Pro Arg Tyr Phe Asp Asn Cys Thr $1650 \hspace{1.5cm} 1660 \hspace{1.5cm}$
- Tyr Arg Asn Leu Ala Ile Pro Met Gly Leu Ala Phe Glu Gly Glu Leu 1665 1670 1680
- Leu Ser Ile Tyr Arg Asn Ser Pro Thr Cys Lys Tyr Gln Val Leu Ser 1700 1705
- 1730 1735 1740

 Leu Tyr Gly Ser Tyr Thr Ile Glu Ala Asp Ala His Thr Leu Ala His
- 1745 1750 1755 176
- Met Met Asn Cys Gly Ala Arg Met Thr Phe 1765 1770
- <210> 445
- <211> 1751
- <212> PRT
- <213> Chlamydia trachomatis serovar D
- <400> 445
- Met Lys Trp Leu Ser Ala Thr Ala Val Phe Ala Ala Val Leu Pro Ser 5 10 15
- Val Ser Gly Phe Cys Phe Pro Glu Pro Lys Glu Leu Asn Phe Ser Arg 20 25 30
- Val Gly Thr Ser Ser Ser Thr Thr Phe Thr Glu Thr Val Gly Glu Ala 35 40 45
- Gly Ala Glu Tyr Ile Val Ser Gly Asn Ala Ser Phe Thr Lys Phe Thr 50 55 60
- Asn Ile Pro Thr Thr Asp Thr Thr Thr Pro Thr Asn Ser Asn Ser Ser 65 70 75 80

Ser Ser Asn Gly Glu Thr Ala Ser Val Ser Glu Asp Ser Asp Ser Thr Thr Thr Pro Asp Pro Lys Gly Gly Gly Ala Phe Tyr Asn Ala His Ser Gly Val Leu Ser Phe Met Thr Arg Ser Gly Thr Glu Gly Ser Leu Thr Leu Ser Glu Ile Lys Ile Thr Gly Glu Gly Gly Ala Ile Phe Ser Gln Gly Glu Leu Leu Phe Thr Asp Leu Thr Gly Leu Thr Ile Gln Asn Asn Leu Ser Gln Leu Ser Gly Gly Ala Ile Phe Gly Glu Ser Thr Ile Ser Leu Ser Gly Ile Thr Lys Ala Thr Phe Ser Ser Asn Ser Ala Glu Val Pro Ala Pro Val Lys Lys Pro Thr Glu Pro Lys Ala Gln Thr Ala Ser Glu Thr Ser Gly Ser Ser Ser Ser Ser Gly Asn Asp Ser Val Ser Ser Pro Ser Ser Ser Arg Ala Glu Pro Ala Ala Ala Asn Leu Gln Ser His Phe Ile Cys Ala Thr Ala Thr Pro Ala Ala Gln Thr Asp Thr Glu Thr Ser Thr Pro Ser His Lys Pro Gly Ser Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Thr Ile Ala Asp Ser Gln Glu Val Leu Phe Ser Ile Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu Lys Asp Val Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly Ala Leu Tyr Val Glu Gly Asp Ile Asn Phe Gln Asp Leu Glu Glu Ile Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr Leu Pro Lys Ala Gln Ala Ser Ala Gly Asn Ala Asp Ala Trp Ala Ser

Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asn Ser Gly Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val Pro Ala Thr Ala Lys Gly Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser Ile Thr Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr Asp Val Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn Ser His Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly Gly Ile Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys Thr Leu Phe Gln Glu Asn Thr Ala Lys Glu Glu Gly Gly Leu Phe Ile Lys Gly Thr Asp Lys Ala Leu Thr Met Thr Gly Leu Asp Ser Phe Cys Leu Ile Asn Asn Thr Ser Glu Lys His Gly Gly Gly Ala Phe Val Thr Lys 545 550 555 560 Glu Ile Ser Gln Thr Tyr Thr Ser Asp Val Glu Thr Ile Pro Gly Ile Thr Pro Val His Gly Glu Thr Val Ile Thr Gly Asn Lys Ser Thr Gly Gly Asn Gly Gly Val Cys Thr Lys Arg Leu Ala Leu Ser Asn Leu Gln Ser Ile Ser Ile Ser Gly Asn Ser Ala Ala Glu Asn Gly Gly Gly Ala His Thr Cys Pro Asp Ser Phe Pro Thr Ala Asp Thr Ala Glu Gln Pro Ala Ala Ala Ser Ala Ala Thr Ser Thr Pro Glu Ser Ala Pro Val Val Ser Thr Ala Leu Ser Thr Pro Ser Ser Ser Thr Val Ser Ser Leu Thr Leu Leu Ala Ala Ser Ser Gln Ala Ser Pro Ala Thr Ser Asn Lys Glu Thr Gln Asp Pro Asn Ala Asp Thr Asp Leu Leu Ile Asp Tyr Val Val Asp Thr Thr Ile Ser Lys Asn Thr Ala Lys Lys Gly Gly Gly Ile 705 710 715 720 Tyr Ala Lys Lys Ala Lys Met Ser Arg Ile Asp Gln Leu Asn Ile Ser

725 730 Glu Asn Ser Ala Thr Glu Ile Gly Gly Gly Ile Cys Cys Lys Glu Ser Leu Glu Leu Asp Ala Leu Val Ser Leu Ser Val Thr Glu Asn Leu Val Gly Lys Glu Gly Gly Leu His Ala Lys Thr Val Asn Ile Ser Asn Leu Lys Ser Gly Phe Ser Phe Ser Asn Asn Lys Ala Asn Ser Ser Ser Thr Gly Val Ala Thr Thr Ala Ser Ala Pro Ala Ala Ala Ala Ala Ser Leu Gln Ala Ala Ala Ala Val Pro Ser Ser Pro Ala Thr Pro Thr Tyr Ser Gly Val Val Gly Gly Ala Ile Tyr Gly Glu Lys Val Thr Phe Ser Gln Cys Ser Gly Thr Cys Gln Phe Ser Gly Asn Gln Ala Ile Asp Asn Asn Pro Ser Gln Ser Ser Leu Asn Val Gln Gly Gly Ala Ile Tyr Ala Lys Thr Ser Leu Ser Ile Gly Ser Ser Asp Ala Gly Thr Ser Tyr Ile Phe Ser Gly Asn Ser Val Ser Thr Gly Lys Ser Gln Thr Thr Gly Gln Ile Ala Gly Gly Ala Ile Tyr Ser Pro Thr Val Thr Leu Asn Cys Pro Ala Thr Phe Ser Asn Asn Thr Ala Ser Met Ala Thr Pro Lys Thr Ser Ser Glu Asp Gly Ser Ser Gly Asn Ser Ile Lys Asp Thr Ile Gly Gly Ala Ile Ala Gly Thr Ala Ile Thr Leu Ser Gly Val Ser Arg Phe Ser Gly Asn Thr Ala Asp Leu Gly Ala Ala Ile Gly Thr Leu Ala Asn Ala Asn Thr Pro Ser Ala Thr Ser Gly Ser Gln Asn Ser Ile Thr Glu Lys Ile Thr Leu Glu Asn Gly Ser Phe Ile Phe Glu Arg Asn Gln Ala Asn Lys Arg Gly Ala Ile Tyr Ser Pro Ser Val Ser Ile Lys Gly Asn 1030 1035 Asn Ile Thr Phe Asn Gln Asn Thr Ser Thr His Asp Gly Ser Ala Ile 1045 1050

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- Tyr Phe Thr Lys Asp Ala Thr Ile Glu Ser Leu Gly Ser Val Leu Phe 1060 1065 1070
- Thr Gly Asn Asn Val Thr Ala Thr Gln Ala Ser Ser Ala Thr Ser Gly 1075
- Gln Asn Thr Asn Thr Ala Asn Tyr Gly Ala Ala Ile Phe Gly Asp Pro 1090 1095 1100
- Gly Thr Thr Gln Ser Ser Gln Thr Asp Ala Ile Leu Thr Leu Leu Ala 1105 111011151115
- Ser Ser Gly Asn Ile Thr Phe Ser Asn Asn Ser Leu Gln Asn Asn Gln 1125 1130 1135
- Gly Asp Thr Pro Ala Ser Lys Phe Cys Ser Ile Ala Gly Tyr Val Lys 1140 1145 1150
- Leu Ser Leu Gln Ala Ala Lys Gly Lys Thr Ile Ser Phe Phe Asp Cys 1155 1160 1165
- Val His Thr Ser Thr Lys Lys Ile Gly Ser Thr Gln Asn Val Tyr Glu 1170 1175 1180
- Thr Leu Asp Ile Asn Lys Glu Glu Asn Ser Asn Pro Tyr Thr Gly Thr 1185 1190 1195 1200
- Ile Val Phe Ser Ser Glu Leu Eis Glu Asn Lys Ser Tyr Ile Pro Gln 1205 1210 1215 Asn Ala Ile Leu Eis Asn Gly Thr Leu Val Leu Lys Glu Lys Thr Glu
- 1220 1225 1230 Leu His Val Val Ser Phe Glu Gln Lys Glu Gly Ser Lys Leu Ile Met
- 1235 1240 1245
 Lys Pro Gly Ala Val Leu Ser Asn Gln Asn Ile Ala Asn Gly Ala Leu

1255

- Val Ile Asn Gly Leu Thr Ile Asp Leu Ser Ser Met Gly Thr Pro Gln
- Ala Gly Glu Ile Phe Ser Pro Pro Glu Leu Arg Ile Val Ala Thr Thr
- 1285 1290 1295 Ser Ser Ala Sér Gly Gly Ser Gly Val Ser Ser Ser Ile Pro Thr Asn
- 1300 1305 1310
 Fro Lys Arg Ile Ser Ala Ala Ala Pro Ser Gly Ser Ala Ala Thr Thr
- Pro Thr Met Ser Glu Asn Lys Val Phe Leu Thr Gly Asp Leu Thr Leu 1330 1340
- Ile Asp Pro Asn Gly Asn Phe Tyr Gln Asn Pro Met Leu Gly Ser Asp 1345 1350 1355 1360
- Leu Asp Val Pro Leu Ile Lys Leu Pro Thr Asn Thr Ser Asp Val Gln 1365 1370 1375

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Val Tyr Asp Leu Thr Leu Ser Gly Asp Leu Phe Pro Gln Lys Gly Tyr 1380 1385 1390

Met Gly Thr Trp Thr Leu Asp Ser Asn Pro Gln Thr Gly Lys Leu Gln 1395 $$1400\$

Asp Asn His Phe Tyr Ala Asn Ser Ile Leu Gly Ser Gln Asn Ser Met 1425 1430 1435

Ile Val Val Lys Gln Gly Leu Ile Asn Asn Met Leu Asn Asn Ala Arg 1445 1450

Phe Asp Asp Ile Ala Tyr Asn Asn Phe Trp Val Ser Gly Val Gly Thr $1460 \hspace{1cm} 1465 \hspace{1cm} 1470 \hspace{1cm}$

Phe Leu Ala Gln Gln Gly Thr Pro Leu Ser Glu Glu Phe Ser Tyr Tyr 1475 1480 1485

Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Lys Pro Arg Gln Asp Phe $1490 \hspace{1.5cm} 1495 \hspace{1.5cm} 1500$

Ile Leu Gly Ala Ala Phe Ser Lys Met Val Gly Lys Thr Lys Ala Ile 1505 \$1510\$

Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Asn Lys Gln 1540 . 1545

His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly 1555 1560 1565

His Ile Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1570 1575 1580

Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 \$1590\$

Ser Met Asp Leu Lys Glu Pro Ser Lys Asp Ser Ser Lys Arg Ile Thr $1605 \hspace{1.5cm} 1610 \hspace{1.5cm} 1615$

Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe Thr 1620 1625 1630

Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Asn 1635 1640 1645

Leu Ser Leu Pro Val Gly Cys Ala Val Glu Gly Ala Ile Met Asn Cys 1650 1655 1660

Asn Ile Leu Met Tyr Asn Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile

Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1685 1690 1695

Ala Gly Gln Val Ile Cys Gly Val Pro Thr Arg Thr Ser Ala Arg Ala

1705

255

1700

Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1720 Asn Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr Ser Cys Gly Ala Arg Met Ile Phe 1745 <210> 446 <211> 660 <212> PRT <213> Chlamydia trachomatis serovar D Met Ser Glu Lys Arg Lys Ser Asn Lys Ile Ile Gly Ile Asp Leu Gly
5 10 15 Thr Thr Asn Ser Cys Val Ser Val Met Glu Gly Gly Gln Pro Lys Val Ile Ala Ser Ser Glu Gly Thr Arg Thr Thr Pro Ser Ile Val Ala Phe 35 45 Lys Gly Glu Thr Leu Val Gly Ile Pro Ala Lys Arg Gln Ala Val Thr Asn Pro Glu Lys Thr Leu Ala Ser Thr Lys Arg Phe Ile Gly Arg Lys Phe Ser Glu Val Glu Ser Glu Ile Lys Thr Val Pro Tyr Lys Val Ala Pro Asn Ser Lys Gly Asp Ala Val Phe Asp Val Glu Gln Lys Leu Tyr Thr Pro Glu Glu Ile Gly Ala Gln Ile Leu Met Lys Met Lys Glu Thr Ala Glu Ala Tyr Leu Gly Glu Thr Val Thr Glu Ala Val Ile Thr 130 135 140 Val Pro Ala Tyr Phe Asn Asp Ser Gln Arg Ala Ser Thr Lys Asp Ala Gly Arg Ile Ala Gly Leu Asp Val Lys Arg Ile Ile Pro Glu Pro Thr 165 170 175 Ala Ala Ala Leu Ala Tyr Gly Ile Asp Lys Glu Gly Asp Lys Lys Ile Ala Val Phe Asp Leu Gly Gly Gly Thr Phe Asp Ile Ser Ile Leu Glu Ile Gly Asp Gly Val Phe Glu Val Leu Ser Thr Asn Gly Asp Thr His 210 215 220 Leu Gly Gly Asp Asp Phe Asp Gly Val Ile Ile Asn Trp Met Leu Asp

225					230					235					240
Glu	Phe	Lys	Lys	Gln 245	Glu	Gly	Ile	Asp	Leu 250	Ser	Lys	Asp	Asn	Met 255	Ala
Leu	Gln	Arg	Leu 260	Lys	Asp	Ala	Ala	Glu 265	Lys	Ala	Lys	Ile	Glu 270	Leu	Ser
Gly	Val	Ser 275	Ser	Thr	Glu	Ile	Asn 280	Gln	Pro	Phe	Ile	Thr 285	Ile	Asp	Ala
Asn	Gly 290	Pro	Lys	His	Leu	Ala 295	Leu	Thr	Leu	Thr	Arg 300	Ala	Gln	Phe	Glu
His 305	Leu	Ala	Ser	Ser	Leu 310	Tle	Glu	Arg	Thr	Lys 315	Gln	Pro	Cys	Ala	Gln 320
Ala	Leu	Lys	Asp	Ala 325	Lys	Leu	Ser	Ala	Ser 330	Asp	Ile	Asp	Asp	Val 335	Leu
Leu	Val	Gly	Gly 340	Met	Ser	Arg	Met	Pro 345	Ala	Val	Gln	Ala	Val 350	Val	Lys
Glu	Ile	Phe 355	Gly	Lys	Glu	Pro	Asn 360	Lys	Gly	Val	Asn	Pro 365	Asp	Glu	Val
Val	Ala 370	Ile	Gly	Ala	Ala	11e 375	Gln	Gly	Gly	Val	Leu 380	Gly	Gly	Glu	Val
Lys 385	Asp	Val	Leu	Leu	Leu 390	Asp	Val	Ile	Pro	Leu 395	Ser	Leu	Gly	Ile	Glu 400
Thr	Leu	Gly	Gly	Val 405	Met	Thr	Pro	Leu	Val 410	Glu	Arg	Asn	Thr	Thr 415	Ile
Pro	Thr	Gln	Lys 420	Lys	Gln	Ile	Phe	Ser 425	Thr	Ala	Ala	Asp	Asn 430	Gln	Pro
Ala	Val	Thr 435	Ile	Val	Val	Leu	Gln 440	Gly	Glu	Arg	Pro	Met 445	Ala	Lys	Asp
Asn	Lys 450	Glu	Ile	Gly	Arg	Phe 455	Asp	Leu	Thr	Asp	Ile 460	Pro	Pro	Ala	Pro
Arg 465	Gly	His	Pro	Gln	Ile 470	Glu	Val	Thr	Phe	Asp 475	Ile	Asp	Ala	Asn	Gly 480
Ile	Leu	His	Val	Ser 485	Ala	Lys	Asp	Ala	Ala 490	Ser	Gly	Arg	Glu	Gln 495	Lys
Ile	Arg	Ile	Glu 500	Ala	Ser	Ser	Gly	Leu 505	Lys	Glu	Asp	Glu	Ile 510	Gln	Gln
Met	Ile	Arg 515	Asp	Ala	Glu	Leu	His 520	Lys	Glu	Glu	Asp	Lys 525	Gln	Arg	Lys
Glu	Ala 530	Ser	Asp	Val	Lys	Asn 535	Glu	Ala	Asp	Gly	Met 540	Ile	Phe	Arg	Ala
Glu 545	Lys	Ala	Val	Lys	Asp 550	Tyr	His	Asp	Lys	Ile 555	Pro	Ala	Glu	Leu	Val 560

Lys Glu Ile Glu Glu His Ile Glu Lys Val Arg Gln Ala Ile Lys Glu Asp Ala Ser Thr Thr Ala Ile Lys Ala Ala Ser Asp Glu Leu Ser Thr His Met Gln Lys Ile Gly Glu Ala Met Gln Ala Gln Ser Ala Ser Ala Ala Ala Ser Ser Ala Ala Asn Ala Gln Gly Gly Pro Asn Ile Asn Ser Glu Asp Leu Lys Lys His Ser Phe Ser Thr Arg Pro Pro Ala Gly Gly 625 630 635 Ser Ala Ser Ser Thr Asp Asn Ile Glu Asp Ala Asp Val Glu Ile Val Asp Lys Pro Glu <210> 447 <211> 326 <212> PRT <213> Chlamydia trachomatis serovar D <400> 447 Met Val Ser Gln Thr Val Ser Val Ala Val Thr Gly Gly Thr Gly Gln Ile Ala Tyr Ser Phe Leu Phe Ser Leu Ala His Gly Asp Val Phe Gly 20 25 30Leu Asp Cys Gly Ile Asp Leu Arg Ile Tyr Asp Ile Pro Gly Thr Glu Arg Ala Leu Ser Gly Val Arg Met Glu Leu Asp Asp Gly Ala Phe Pro Leu Leu Gln Arg Val Gln Val Thr Thr Ser Leu His Asp Ala Phe Asp Gly Ile Asp Ala Ala Phe Leu Ile Gly Ser Val Pro Arg Gly Pro Gly Met Glu Arg Arg Asp Leu Leu Lys Lys Asn Gly Glu Ile Phe Ala Thr Gln Gly Lys Ala Leu Asn Thr Thr Ala Lys Arg Asp Ala Lys Ile Phe Val Val Gly Asn Pro Val Asn Thr Asn Cys Trp Ile Ala Met Asn His 135 Ala Pro Arg Leu Leu Arg Lys Asn Phe His Ala Met Leu Arg Leu Asp Gln Asn Arg Met His Ser Met Leu Ser His Arg Ala Glu Val Pro Leu

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Ser Ala Val Ser Gln Val Val Val Trp Gly Asn His Ser Ala Lys Gln 185 Val Pro Asp Phe Thr Gln Ala Leu Ile Asn Asp Arg Pro Ile Ala Glu Thr Ile Ala Asp Arg Asp Trp Leu Glu Asn Ile Met Val Pro Ser Val Gln Ser Arg Gly Ser Ala Val Ile Glu Ala Arg Gly Lys Ser Ser Ala Ala Ser Ala Ala Arg Ala Leu Ala Glu Ala Ala Arg Ser Ile Tyr Gln 245 250 255 Pro Lys Glu Gly Glu Trp Phe Ser Ser Gly Val Cys Ser Asp His Asn 260 265 270 Pro Tyr Gly Leu Pro Glu Asp Leu Ile Phe Gly Phe Pro Cys Arg Met Leu Ala Thr Gly Glu Tyr Glu Val Ile Pro Arg Leu Pro Trp Asp Ala Phe Ile Arg Gly Lys Met Gln Ile Ser Leu Asp Glu Ile Leu Gln Glu Lys Ala Ser Val Ser Leu <210> 448 <211> 232 <212> PRT <213> Chlamydia trachomatis serovar D <400> 448 Met Thr Lys His Gly Lys Arg Ile Arg Gly Ile Gln Glu Thr Tyr Asp Leu Ala Lys Ser Tyr Ser Leu Gly Glu Ala Ile Asp Ile Leu Lys Gln Cys Pro Thr Val Arg Phe Asp Gln Thr Val Asp Val Ser Val Lys Leu

Cys Pro 1 Arg Phe Asp Gln Thr Val Asp Val Ser Val Lys Leu 45 Val Leu Arg Ile Asp Pro Arg Lys Ser Asp Gln Gln Ile Arg Gly Ser Val Ser Ser For His Gly Thr Gly Lys Val Leu Arg Ile Arg Clu Val Phe Ala Ala 65 For Asp Lys Ala Ala Glu Ala Ile Glu Ala Gly Ala Asp Phe Val Gly 90 Val Asp Lys Ala Ala Glu Lys Ile Lys Gly Gly Trp Val Asp Phe Asp 100 Val Ala Val Ala Thr Fro Asp Met Arg Glu Val Gly Lys Leu Gly 115 Val Ala Val Ala Thr Fro Asp Met Met Arg Glu Val Gly Lys Leu Gly 115 Val Ala Val Ala Thr Fro Asp Met Met Arg Glu Val Gly Lys Leu Gly 115 Val Ala Val Ala Thr Fro Asp Met Met Arg Glu Val Gly Lys Leu Gly 115 Val Ala Val Ala Thr Fro Asp Met Met Arg Glu Val Gly Lys Leu Gly 115 Val Asp Phe Asp 110 Val Asp Val Val Gly Lys Leu Gly 115 Val Asp Val Val Gly Lys Leu Gly 115 Val Asp Val Val Gly Val Gly Val Gly Val Gly Val Gly 115 Val Asp Val Val Gly
Lys Val Leu Gly Pro Arg Asn Leu Met Pro Thr Pro Lys Ala Gly Thr 135 Val Thr Thr Asp Val Val Lys Thr Val Ala Glu Leu Arg Lys Gly Lys Ile Glu Phe Lys Ala Asp Arg Ala Gly Val Cys Asn Val Gly Val Ala Lys Leu Ser Phe Asp Ser Ala Gln Ile Lys Glu Asn Val Glu Ala Leu Cys Ala Ala Leu Val Lys Ala Lys Pro Ala Thr Ala Lys Gly Gln Tyr Leu Val Asn Phe Thr Ile Ser Ser Thr Met Gly Pro Gly Val Thr Val 210 215 220 Asp Thr Arg Glu Leu Ile Ala Leu <210> 449 <211> 1252 <212> PRT <213> Chlamydia trachomatis serovar D <400> 449 Met Phe Lys Cys Pro Glu Arg Val Ser Ile Lys Lys Lys Glu Asp Ile Leu Asp Leu Pro Asn Leu Val Glu Val Gln Ile Lys Ser Tyr Lys Gln Phe Leu Gln Ile Gly Lys Leu Ala Glu Glu Arg Glu Asn Ile Gly Leu Glu Glu Val Phe Arg Glu Ile Phe Pro Ile Lys Ser Tyr Asn Glu Ala Thr Ile Leu Glu Tyr Leu Ser Tyr Asn Leu Gly Val Pro Lys Tyr Ser Pro Glu Glu Cys Ile Arg Arg Gly Ile Thr Tyr Ser Val Thr Leu Lys Val Arg Phe Arg Leu Thr Asp Glu Thr Gly Ile Lys Glu Glu Glu Val Tyr Met Gly Thr Ile Pro Ile Met Thr Asp Lys Gly Thr Phe Ile Ile Asn Gly Ala Glu Arg Val Val Val Ser Gln Val His Arg Ser Pro Gly Ile Asn Phe Glu Gln Glu Lys His Ser Lys Gly Asn Val Leu Phe Ser Phe Arg Ile Ile Pro Tyr Arg Gly Ser Trp Leu Glu Ala Val Phe Asp

Ile	Asn	Asp	Leu 180	Ile	Tyr	Ile	His	11e 185	Asp	Arg	Lys	Lys	Arg 190	Arg	Arg
Lys	Ile	Leu 195	Ala	Met	Thr	Phe	11e 200	Arg	Ala	Leu	Gly	Tyr 205	Ser	Thr	Asp
Ala	Asp 210	Ile	Ile	Glu	Glu	Phe 215	Phe	Ser	Val	Glu	Glu 220	Arg	Ser	Leu	Arg
Leu 225	Glu	Lys	Asp	Phe	Val 230	Ala	Leu	Val	Gly	Lys 235	Val	Leu	Ala	Asp	Asn 240
Val	Val	Asp	Ala	Asp 245	Ser	Ser	Leu	Val	Tyr 250	Gly	Lys	Ala	Gly	Glu 255	Lys
Leu	Ser	Thr	Ala 260	Met	Leu	Lys	Arg	11e 265	Leu	Asp	Ala	Gly	Val 270	Gln	Ser
Leu	Lys	Ile 275	Ala	Val	Gly	Ala	Asp 280	Glu	Asn	His	Pro	11e 285	Ile	Ьуз	Met
Leu	Ala 290	Lys	Asp	Pro	Thr	Asp 295	Ser	Tyr	Glu	Ala	Ala 300	Leu	Lys	Asp	Phe
Tyr 305	Arg	Arg	. Leu	Arg	Pro 310	Gly	Glu	Pro	Ala	Thr 315	Leu	Val	Asn	Ala	Arg 320
Ser	Thr	Ile	Met	Arg 325	Leu	Phe	Phe	Asp	Ala 330	Lys	Arg	Tyr	Asn	Leu 335	Gly
Arg	Va1	Gly	Arg 340	Tyr	Lys	Leu	Asn	Lys 345	Lys	Leu	Gly	Phe	Pro 350	Leu	Asp
Asp	Glu	Thr 355	Leu	Ser	Gln	Val	Thr 360	Leu	Arg	Lys	Glu	Asp 365	Val	Ile	Gly
Ala	-														
	370	Lys	Tyr	Leu	Ile	Arg 375	Leu	Arg	Met	Gly	Asp 380	Glu	Lys	Thr	Ser
Ile 385	370					375					380			Thr Val	
385	370 Asp	Asp	Ile	Asp	His 390	375 Leu	Ala	Asn	Arg	Arg 395	380 Val	Arg	Ser		Gly 400
385 Glu	370 Asp Leu	Asp Ile	Ile Gln	Asp Asn 405	His 390 His	375 Leu Cys	Ala Arg	Asn Ser	Arg Gly 410	Arg 395 Leu	380 Val Ala	Arg Arg	Ser Met	Val Glu	Gly 400 Lys
385 Glu Ile	Asp Leu Val	Asp Ile Arg	Ile Gln Glu 420	Asp Asn 405 Arg	His 390 His Met	375 Leu Cys Asn	Ala Arg Leu	Asn Ser Phe 425	Arg Gly 410 Asp	Arg 395 Leu Phe	Val Ala Ser	Arg Arg Ser	Ser Met Asp 430	Val Glu 415	Gly 400 Lys Leu
385 Glu Ile Thr	370 Asp Leu Val Pro	Asp Ile Arg Gly 435	Ile Gln Glu 420 Lys	Asp Asn 405 Arg	His 390 His Met	375 Leu Cys Asn Ser	Ala Arg Leu Ala 440	Asn Ser Phe 425 Lys	Arg Gly 410 Asp Gly	Arg 395 Leu Phe Leu	380 Val Ala Ser Val	Arg Arg Ser Ser 445	Ser Met Asp 430 Val	Val Glu 415 Thr	Gly 400 Lys Leu Lys
385 Glu Ile Thr	370 Asp Leu Val Pro Phe 450	Asp Ile Arg Gly 435 Phe	Ile Gln Glu 420 Lys Ser	Asp Asn 405 Arg Ile Arg	His 390 His Met Ile Ser	375 Leu Cys Asn Ser Gln 455	Ala Arg Leu Ala 440 Leu	Asn Ser Phe 425 Lys Ser	Arg Gly 410 Asp Gly Gln	Arg 395 Leu Phe Leu	380 Val Ala Ser Val Met 460	Arg Arg Ser Ser 445 Asp	Ser Met Asp 430 Val	Val Glu 415 Thr	Gly 400 Lys Leu Lys Asn

Ala	Ser	His	Tyr 500	Gly	Arg	Ile	Cys	Pro 505	Ile	Glu	Thr	Pro	Glu 510	Gly	Pro
Asn	Ile	Gly 515	Leu	Ile	Thr	Ser	Leu 520	Ser	Ser	Phe	Ala	Lys 525	Ile	Asn	G1u
Phe	Gly 530	Phe	Ile	Glu	Thr	Pro 535	Tyr	Arg	Val	Val	Arg 540	Asp	Gly	Ile	Val
Thr 545	Asp	Glu	Ile	Glu	Tyr 550	Met	Thr	Ala	Asp	Val 555	G1u	Glu	Glu	Cys	Val 560
Ile	Ala	G1n	Ala	Ser 565	Ala	Glu	Leu	Asp	Glu 570	Tyr	Asp	Met	Phe	Lys 575	Thr
Pro	Val	Cys	Trp 580	Ala	Arg	Tyr	Lys	Gly 585	Glu	Ala	Phe	Glu	Ala 590	Asp	Thr
Ser	Thr	Val 595	Thr	His	Met	Asp	Val 600	Ser	Pro	Lys	Gln	Leu 605	Val	Ser	Val
Val	Thr 610	Gly	Leu	Ile	Pro	Phe 615	Leu	Glu	His		Asp 620	Ala	Asn	Arg	Ala
Leu 625	Met	Gly	Ser	Asn	Met 630	Gln	Arg	Gln	Ala	Val 635	Pro	Leu	Leu	Lys	Thr 640
Glu	Ala	Ala	Ile	Val 645	Gly	Thr	Gly	Leu	Glu 650	Gly	Arg	Ala	Ala	Lys 655	Asp
Ser	Gly	Ala	11e 660	Ile	Val	Ala	Gln	Glu 665	Asp	Gly	Val	Val	Glu 670	Tyr	Val
Asp	Ser	Tyr 675	Glu	Ile	Val	Val	Ala 680	Lys	Lys	Asn	Asn	Pro 685	Thr	Leu	Lys
Asp	Arg 690	Tyr	Gln	Leu	Lys	Lys 695	Phe	Leu	Arg	Ser	Asn 700	Ser	Gly	Thr	Cys
Ile 705	Asn	Gln	Thr	Pro	Leu 710	Cys	Ser	Val	Gly	Asp 715	Val	Val	Thr	His	Gly 720
Asp	Val	Leu	Ala	Asp 725	Gly	Pro	Ala	Thr	Asp 730	Lys	Gly	Glu	Leu	Ala 735	Leu
Gly	Lys	Asn	Val 740	Leu	Val	Ala	Phe	Met 745	Pro	Trp	Tyr	Gly	Tyr 750	Asn	Phe
Glu	Asp	Ala 755	Ile	Ile	Ile	Ser	Glu 760	Arg	Leu	Ile	Lys	Gln 765	Asp	Ala	Tyr
Thr	Ser 770	Ile	Tyr	Ile	Glu	Glu 775	Phe	Glu	Leu	Thr	Ala 780	Arg	Asp	Thr	Lys
Leu 785	Gly	Lys	G1u	Glu	11e 790	Thr	Arg	Asp	Ile	Pro 795	Asn	Val	Ser	Glu	Glu 800
Val	Leu	Ala	Asn	Leu 805	Gly	Glu	Asp	Gly	Val 810	Val	Arg	Ile	Gly	Ala 815	Glu
Val	Lys	Pro	Gly	Asp	Ile	Leu	Val	Gly	Lys	Ile	Thr	Pro	Lys	Ser	Glu

			820					825					830		
Thr	Glu	Leu 835	Ala	Pro	Glu	Glu	Arg 840	Leu	Leu	Arg	Ala	11e 845	Phe	Gly	Glu
Lys	Ala 850	Ala	Asp	Val	Lys	Asp 855	Ala	Ser	Leu	Thr	Val 860	Pro	Pro	Gly	Thr
Glu 865	Gly	Val	Val	Met	Asp 870	Val	Lys	Val	Phe	Ser 875	Arg	Lys	Asp	Arg	Leu 880
Ser	Lys	Ser	Asp	Asp 885	Glu	Leu	Val	Glu	Glu 890	Ala	Val	His	Leu	Lys 895	Asp
Leu	Gln	Lys	Glu 900	Tyr	Lys	Ser	Gln	Leu 905	Ala	Gln	Leu	Lys	Val 910	Glu	His
Arg	Glu	Lys 915	Leu	Gly	Ala	Leu	Leu 920	Leu	Asn	Glu	Lys	Ala 925	Pro	Ala	Ala
Ile	Ile 930	His	Arg	Arg	Ser	Ala 935	Asp	Ile	Leu	Val	Gln 940	Glu	Gly	Ala	Ile
Phe 945	Asp	Gln	Glu	Thr	Ile 950	Glu	Leu	Leu	Glu	Arg 955	Glu	Ser	Leu	Val	Asp 960
Leu	Leu	Met	Ala	Pro 965	Cys	Asp	Met	Tyr	Asp 970	Val	Leu	Lys	Asp	11e 975	Leu
Ser	Ser	Tyr	Glu 980	Thr	Ala	Val	Gln	Arg 985	Leu	Glu	Val	Asn	Tyr 990	Lys	Thr
Glu	Ala	Glu 995	His	Ile	Lys	Glu	Gly 1000		Ala	Asp	Leu	Asp 1005		Gly	Val
Ile	Arg 1010		Val	Lys	Val	Tyr 1015		Ala	Ser	Lys	Arg 1020		Leu	Gln	Val
Gly 1025		Lys	Met	Ala	Gly 1030		His	Gly	Asn	Lys 1035		Val	Val	Ser	Lys 1040
Ile	Val	Pro	Glu	Ala 1045		Met	Pro	Phe	Leu 1050	Ala	Asn	Gly	Glu	Thr 1055	
Gln	Met	Ile	Leu 1060	Asn)	Pro	Leu	Gly	Va1 1065	Pro	Ser	Arg	Met	Asn 1070		Gly
Gln	Val	Leu 1075		Thr	His	Leu	Gly 1080		Ala	Ala	Lys	Thr 1085		Gly	Ile
Tyr	Val 1090		Thr	Pro	Val	Phe 1095		Gly	Phe	Pro	Glu 1100		Arg	Ile	Trp
Asp 1105	Met	Met	Ile	Glu	Gln 1110		Leu	Pro	Glu	Asp 1115		Lys	Ser	Tyr	Leu 1120
Phe	Asp	Gly	Lys	Thr 1125		Glu	Arg	Phe	Asp 1130	Ser	Lys	Val	Val	Val 1135	
Tyr	Ile	Tyr	Met 1140		Lys	Leu	Ser	His 1145		Ile	Ala	Asp	Lys 1150		His

Ala Arg Ser Ile Gly Pro Tyr Ser Leu Val Thr Gln Gln Pro Leu Gly 1160 Gly Lys Ala Gln Met Gly Gly Gln Arg Phe Gly Glu Met Glu Val Trp Ala Leu Glu Ala Tyr Gly Val Ala His Met Leu Gln Glu Ile Leu Thr Val Lys Ser Asp Asp Val Ser Gly Arg Thr Arg Ile Tyr Glu Ser Ile Val Lys Gly Glu Asn Leu Leu Arg Ser Gly Thr Pro Glu Ser Phe Asn Val Leu Ile Lys Glu Met Gln Gly Leu Gly Leu Asp Val Arg Pro Met 1235 1240 1245 Val Val Asp Ala 1250 <210> 450 <211> 298 <212> PRT <213> Chlamydia trachomatis serovar D <400> 450 Met Leu Lys Ile Asp Leu Thr Gly Lys Ile Ala Phe Ile Ala Gly Ile Gly Asp Asp Asn Gly Tyr Gly Trp Gly Ile Ala Lys Met Leu Ala Glu 20 25 30 Ala Gly Ala Thr Ile Leu Val Gly Thr Trp Val Pro Ile Tyr Lys Ile Phe Ser Gln Ser Leu Glu Leu Gly Lys Phe Asn Ala Ser Arg Glu Leu Ser Asn Gly Glu Leu Leu Thr Phe Ala Lys Ile Tyr Pro Met Asp Ala 65 70 75 80 Ser Phe Asp Thr Pro Glu Asp Ile Pro Gln Glu Ile Leu Glu Asn Lys Arg Tyr Lys Asp Leu Ser Gly Tyr Thr Val Ser Glu Val Val Glu Gln Val Lys Lys His Phe Gly His Ile Asp Ile Leu Val His Ser Leu Ala Asn Ser Pro Glu Ile Ala Lys Pro Leu Leu Asp Thr Ser Arg Lys Gly Tyr Leu Ala Ala Leu Ser Thr Ser Ser Tyr Ser Phe Ile Ser Leu Leu Ser His Phe Gly Pro Ile Met Asn Ala Gly Ala Ser Thr Ile Ser Leu

Thr Tyr Leu Ala Ser Met Arg Ala Val Pro Gly Tyr Gly Gly Gly Met Asn Ala Ala Lys Ala Ala Leu Glu Ser Asp Thr Lys Val Leu Ala Trp Glu Ala Gly Arg Arg Trp Gly Val Arg Val Asn Thr Ile Ser Ala Gly Pro Leu Ala Ser Arg Ala Gly Lys Ala Ile Gly Phe Ile Glu Arg Met Val Asp Tyr Tyr Gln Asp Trp Ala Pro Leu Pro Ser Pro Met Glu Ala Glu Gln Val Gly Ala Ala Ala Phe Leu Val Ser Pro Leu Ala Ser Ala Ile Thr Gly Glu Thr Leu Tyr Val Asp His Gly Ala Asn Val Met Gly Ile Gly Pro Glu Met Phe Pro Lys Asp 295 <210> 451 <211> 298 <212> PRT <213> Chlamydia trachomatis serovar D <400> 451 Met Ser Leu Gln Lys Leu Leu Val Thr Asp Ile Asp Gly Thr Ile Thr His Gln Ser His Leu Leu His Asp Arg Val Val Lys Ala Leu His Gln Tyr Tyr Asp Ser Gly Trp Gln Leu Phe Phe Leu Thr Gly Arg Tyr Phe Ser Tyr Ala Tyr Pro Leu Phe Gln Asn Phe Ser Val Pro Phe Leu Leu Gly Ser Gln Asn Gly Ser Ser Val Trp Ser Ser Thr Asp Lys Glu Phe Ile Tyr Phe Arg Ser Leu Ser Arg Asp Phe Leu Tyr Val Leu Glu Lys Tyr Phe Glu Asp Leu Asp Leu Ile Ala Cys Ile Glu Ser Gly Ala Ser Asn Arg Asp Val Tyr Phe Arg Lys Gly Leu Gly Lys Thr Ser Gln Glu Leu Lys Ala Ile Leu Asp Ala Val Tyr Phe Pro Thr Pro Glu Ala Ala Arg Leu Leu Val Asp Val Gln Gly His Leu Ser Glu Glu Phe Ser Tyr

Glu Asp Phe Ala Ile Ala Lys Phe Phe Gly Glu Arg Glu Glu Val Lys Lys Ile Met Asp Arg Phe Ile Gln Ser Pro Glu Val Ser Ser Gln Val Thr Met Asn Tyr Met Arg Trp Pro Phe Asp Phe Lys Tyr Ala Val Leu Leu Leu Thr Leu Lys Asp Val Ser Lys Gly Phe Ala Val Asp Gln Val 210 215 220 Val Gln Thr Phe Tyr Lys Glu Asn Lys Pro Phe Ile Met Ala Ser Gly Asp Asp Ala Asn Asp Ile Asp Leu Leu Ser Arg Gly Asp Phe Lys Ile Val Ile Gln Thr Ala Pro Glu Glu Met His Gly Leu Ala Asp Phe Leu Ala Pro Pro Ala Lys Asp Phe Gly Ile Leu Ser Ala Trp Glu Ala Gly 275 280 285 Glu Leu Arg Tyr Lys Gln Leu Val Asn Pro <210> 452 <211> 153 <212> PRT <213> Chlamydia trachomatis serovar D <400> 452 Met Leu Arg Leu Phe Gln His Ile Leu Cys Phe Leu Glu Glu Asp Pro Ser Phe Val Asp Val Pro Gln Glu Leu Ser Phe Val Asn Glu Ala Phe 20 30Ser Gly Ser Met Arg Trp Glu Val Gly Arg Met Leu Gly Ser Leu Leu 35 40 45 Leu Leu Gly Ile Phe Gly Gly Gly Cys Leu Leu Phe Arg Arg Phe 50 60Leu Arg Ser Arg Gly His Leu Pro Ser Gly Asn Ser Ser Ile Lys Ile Leu Asp Gln Arg Val Leu Ala Ser Lys Thr Ser Ile Tyr Val Ile Lys Val Ala Asn Lys Thr Leu Val Val Ala Glu Arg Gly Glu Arg Val Thr Leu Leu Ser Glu Phe Pro Pro Asn Thr Asp Leu Asn Glu Leu Ile Gln 120 Lys Asp Gln Lys Lys Pro Ser Thr Pro Arg Gly Glu Met Leu Ser Gly

Phe Leu Lys Gln Phe Lys Glu Lys Lys <210> 453 <211> 569 <212> PRT <213> Chlamvdia trachomatis serovar D Met Pro Lys Gln Ala Asp Tyr Thr Trp Gly Ala Lys Lys Asn Leu Asp Thr Ile Ala Cys Leu Pro Glu Asp Val Lys Gln Phe Lys Asp Leu Leu 20 25 30 Tyr Ala Met Tyr Gly Phe Thr Ala Thr Glu Glu Glu Pro Thr Ser Glu Val His Pro Gly Ala Ile Leu Lys Gly Thr Val Val Asp Ile Ser Lys Asp Phe Val Val Val Asp Val Gly Leu Lys Ser Glu Gly Val Ile Pro Met Ser Glu Phe Ile Asp Ser Ser Glu Gly Leu Thr Val Gly Ala Glu 85 90 95 Val Glu Val Tyr Leu Asp Gln Thr Glu Asp Asp Glu Gly Lys Val Val 100 105 Leu Ser Arg Glu Lys Ala Thr Arg Gln Arg Gln Trp Glu Tyr Ile Leu 115 120 125 Ala His Cys Glu Glu Gly Ser Ile Val Lys Gly Gln Ile Thr Arg Lys Val Lys Gly Gly Leu Ile Val Asp Ile Gly Met Glu Ala Phe Leu Pro Gly Ser Gln Ile Asp Asn Lys Lys Ile Lys Asn Leu Asp Asp Tyr Val 165 170 175Gly Lys Val Cys Glu Phe Lys Ile Leu Lys Ile Asn Val Asp Arg Arg Asn Val Val Ser Arg Arg Glu Leu Leu Glu Ala Glu Arg Ile Ser Lys Lys Ala Glu Leu Ile Glu Gln Ile Thr Ile Gly Glu Arg Arg Lys $210 \hspace{1.5cm} 225 \hspace{1.5cm} 220 \hspace{1.5cm}$ Gly Ile Val Lys Asn Ile Thr Asp Phe Gly Val Phe Leu Asp Leu Asp Gly Ile Asp Gly Leu Leu His Ile Thr Asp Met Thr Trp Lys Arg Ile Arg His Pro Ser Glu Met Val Glu Leu Asn Gln Glu Leu Glu Val Ile

Ile	Leu	Ser 275	Val	Asp	Lys	Glu	Lys 280	Gly	Arg	Val	Ala	Leu 285	Gly	Leu	Lys
Gln	Lys 290	Glu	His	Asn	Pro	Trp 295	Glu	Asp	Ile	Glu	Lys 300	Lys	Tyr	Pro	Pro
Gly 305	Lys	Arg	Val	Arg	Gly 310	Lys	Ile	Val	Lys	Leu 315	Leu	Pro	Tyr	Gly	Ala 320
Phe	Ile	Glu	Ile	Glu 325	Glu	Gly	Ile	Glu	Gly 330	Leu	Ile	His	Val	Ser 335	Glu
Met	Ser	Trp	Val 340	Lys	Asn	Ile	Val	Asp 345	Pro	Asn	Glu	Val	Val 350	Asn	Lys
Gly	Asp	Glu 355	Val	Glu	Val	Val	Val 360	Leu	Ser	Ile	Gln	Lys 365	Asp	Glu	Gly
Lys	Ile 370	Ser	Leu	Gly	Leu	Lys 375	Gln	Thr	Lys	His	Asn 380	Pro	Trp	Asp	Asn
Ile 385	Glu	G1u	Lys	Tyr	Pro 390	Ile	Gly	Leu	Arg	Val 395	Thr	Ala	Glu	Ile	Lys 400
Asn	Leu	Thr	Asn	Tyr 405	Gly	Ala	Phe	Val	Glu 410	Leu	Glu	Pro	Gly	Ile 415	Glu
Gly	Leu	Ile	His 420	Ile	Ser	Asp	Met	Ser 425	Trp	Ile	Lys	Lys	Val 430	Ser	His
Pro	Ser	Glu 435	Leu	Phe	Lys	Lys	Gly 440	Asn	Thr	Val	Glu	Ala 445	Val	Ile	Leu
Ser	Val 450	Asp	Lys	Glu	Ser	Lys 455	Lys	Ile	Thr	Leu	Gly 460	Val	Lys	Gln	Leu
Thr 465	Pro	Asn	Pro	Trp	Asp 470	Glu	Ile	Glu	Val	Met 475	Phe	Pro	Val	Gly	Ser 480
Asp	Ile	Ser	Gly	Val 485	Val	Thr	Lys	Ile	Thr 490	Ala	Phe	Gly	Ala	Phe 495	Val
Glu	Leu	Gln	Asn 500	Gly	Ile	Glu	Gly	Leu 505	Ile	His	Val	Ser	Glu 510	Leu	Ser
Glu	Lys	Pro 515	Phe	Ala	Lys	Ile	Glu 520	Asp	Val	Leu	Ser	Ile 525	Gly	Asp	Lys
Val	Ser 530	Ala	Lys	Val	Ile	Lys 535	Leu	Asp	Pro	Asp	His 540	Lys	Lys	Val	Ser
Leu 545	Ser	Ile	Lys	Glu	Phe 550	Leu	Va1	His	Gly	Gly 555	Asp	Ala	Gly	His	Asp 560
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Thr 305	Thr	Leu	Lys	Trp	Ile 310	Leu	Asp	Ala	Leu	Arg 315	Tyr	Trp	Val	Gln	Glu 320
Met	His	Val	Asp	Gly 325	Phe	Arg	Phe	Asp	Leu 330	Ala	Ser	Val	Phe	Ser 335	Arg
Asp	Pro	Gln	Gly 340	Val	Pro	Leu	Pro	Leu 345	Thr	Pro	Ile	Leu	Gln 350	Ala	Ile
Ser	Ser	Asp 355	Ser	Ile	Leu	Ser	Glu 360	Thr	Lys	Leu	Ile	Ala 365	Glu	Pro	Trp
Asp	Ala 370	Gly	Gly	Leu	Tyr	Gln 375	Leu	Gly	His	Phe	Pro 380	Ser	Ile	Ser	Thr
Arg 385	Trp	Ser	Glu	Trp	Asn 390	Gly	Cys	Tyr	Arg	Asp 395	His	Val	Lys	Ala	Phe 400
Leu	Asn	Gly	Asp	Ala 405	His	Gln	Val	Ser	Ser 410	Phe	Ala	Ser	Arg	Ile 415	Ser
Gly	Ser	His	Asp 420	Ile	Tyr	Pro	Asn	Gly 425	Lys	Pro	Thr	Asn	Ser 430	Ile	Asn
Tyr	Ile	Cys 435	Ser	His	Asp	Gly	Phe 440	Thr	Leu	Tyr	Asp	Thr 445	Val	Ala	Tyr
Asn	Asp 450	Lys	His	Asn	Glu	Glu 455	Asn	Gly	Glu	Tyr	Asn 460	Arg	Asp	Gly	Thr
Ser 465	Ala	Asn	Tyr	Ser	Tyr 470	Asn	Phe	Gly	Cys	Glu 475	Gly	Glu	Thr	Thr	Asp 480
Pro	Thr	Ile	Cys	Ala 485	Leu	Arg	Glu	Arg	Gln 490	Met	Lys	Asn	Phe	Phe 495	Leu
Ala	Leu	Phe	Leu 500	Ser	Gln	Gly	Ile	Pro 505	Met	Ile	G1n	Ser	Gly 510	Asp	Glu
Tyr	Gly	His 515	Thr	Ala	Tyr	Gly	Asn 520	Asn ,	Asn	His	Trp	Cys 525	Leu	Asp.	Thr
	11e 530					535					540		_		
Phe 545	Ser	Phe	Leu	Cys	Gln 550	Val	Ile	Ala	Leu	Arg 555	Lys	Ala	Tyr	Thr	G1u 560
Leu	Phe	Asn	Thr	Ser 565	Phe	Leu	Ser	Glu	Asp 570	Thr	Ile	Thr	Trp	Leu 575	Asn
Thr	Lys	Gly	Ser 580	Pro	Arg	Glu	Trp	Gly 585	Ala	Asp	His	Tyr	Leu 590	Ala	Phe
Glu	Leu	Lys 595	His	Leu	Asn	Tyr	Ser 600	Leu	Phe	Val	Ala	Phe 605	Tyr	Ser	Gly
Asn	Glu 610	Arg	Ile	Glu	Ile	Ser 615	Leu	Pro	Lys	Pro	Arg 620	Lys	Glu	His	Leu
Ala	Tyr	Glu	Lys	Ile	Val	Asp	Ser	Thr	Thr	Gly	Phe	Phe	Ser	Gln	Ile

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625	630	635	640
Leu Ser Pro Lys Leu 645	Ser Leu Glu Pro Tyr 650		la Ile 55
Ser Arg Arg Lys Thr 660	Ser Leu Glu Ser Arg 665		
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100

90

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120 125

WO 02/08267 PCT/US01/23121 289

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545 550 555 560
Tyr Gly Gly Asp Ile Thr Arg Lys Arg Lys Leu Trp Glu Lys Gln Lys
565 570 575
Lys Gly Lys Lys Arg Met Lys Glu Phe Gly Lys Val Ser Ile Pro Asn
            580
                           585
Thr Ala Phe Ile Glu Val Leu Lvs Leu Asp
        595
                             600
<210> 496
<211> 324
<212> PRT
<213> Chlamydia pneumoniae
<400> 496
Met Glu Leu Pro His Glu Lys Gln Val Val Glu Tyr Glu Lys Ala
                                  10
Ile Ala Glu Phe Lys Glu Lys Asn Lys Lys Asn Ser Leu Leu Ser Ser
                                 25
Ser Glu Ile Gln Lys Leu Glu Lys Arg Leu Asp Lys Leu Lys Glu Lys 35 40 45
Ile Tyr Ser Asp Leu Thr Pro Trp Glu Arg Val Gln Ile Cys Arg His
                      55
Pro Ser Arg Pro Arg Thr Val Asn Tyr Ile Glu Gly Met Cys Glu Glu
65 70 75 80
Phe Val Glu Leu Cys Gly Asp Arg Thr Phe Arg Asp Asp Pro Ala Val
85 90 95
Val Gly Gly Phe Val Lys Ile Gln Gly Gln Arg Phe Val Leu Ile Gly
100 105 110
                                                       110
Gln Glu Lys Gly Cys Asp Thr Ala Ser Arg Leu His Arg Asn Phe Gly
                             120
Met Leu Cys Pro Glu Gly Phe Arg Lys Ala Leu Arg Leu Gly Lys Leu
                                           140
Ala Glu Lys Phe Gly Leu Pro Val Val Phe Leu Val Asp Thr Pro Gly 145 150 155 160
Ala Tyr Pro Gly Leu Thr Ala Glu Glu Arg Gly Gln Gly Trp Ala Ile
165 170 175
Ala Lys Asn Leu Phe Glu Leu Ser Arg Leu Ala Thr Pro Val Ile Ile
180 185 190
Val Val Ile Gly Glu Gly Cys Ser Gly Gly Ala Leu Gly Met Ala Val
195 200 205
Gly Asp Ser Val Ala Met Leu Glu His Ser Tyr Tyr Ser Val Ile Ser
210 215 220
Pro Glu Gly Cys Ala Ser Ile Leu Trp Lys Asp Pro Lys Lys Asn Ser
                     230
                                         235
Glu Ala Ala Ser Met Leu Lys Met His Gly Glu Asn Leu Lys Gln Phe 245 250 255
Gly Ile Ile Asp Thr Val Ile Lys Glu Pro Ile Gly Gly Ala His His 260 265 270
Asp Pro Ala Leu Val Tyr Ser Asn Val Arg Glu Phe Ile Ile Gln Glu 275 280 285
Trp Leu Arg Leu Lys Asp Leu Ala Ile Glu Glu Leu Leu Glu Lys Arg
290 295 300
Tyr Glu Lys Phe Arg Ser Ile Gly Leu Tyr Glu Thr Thr Ser Glu Ser
                     310
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Gly Pro Glu Ala

<210> 497

<211> 659

<212> PRT

<213> Chlamydia pneumoniae

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145 150 155 160 150 155 Asp Tyr Phe Lys Ala Leu Gln Gln Leu Pro Met Thr Phe Phe His Asp 165 170 175 His Asp Ile Gly Asn Leu Ser Asn Arg Val Met Thr Asp Ser Ala Ser Ile Ala Leu Ala Val Asn Ser Leu Met Ile Asn Tyr Ile Gln Ala Pro 205 200 Ile Thr Phe Ile Leu Thr Leu Gly Val Cys Leu Ser Ile Ser Trp Lys 215 220 Phe Ser Ile Leu Ile Cys Val Ala Phe Pro Ile Phe Ile Leu Pro Ile 230 235 Val Val Ile Ala Arg Lys Ile Lys Asn Leu Ala Lys Arg Ile Gln Lys 245 250 255 Ser Gln Asp Ser Phe Ser Ser Val Leu Tyr Asp Phe Leu Ala Gly Val 265 Met Thr Val Lys Val Phe Arg Thr Glu Lys Phe Ala Phe Thr Lys Tyr 275 280 285 Cys Glu His Asn Asn Lys Ile Ser Ala Leu Glu Glu Lys Ser Ala Ala 290 295 300 Tyr Gly Leu Leu Pro Arg Pro Leu Leu His Thr Ile Ala Ser Leu Phe 305 310 315 320 315 Phe Ala Phe Val Val Val Ile Gly Ile Tyr Lys Phe Ala Ile Pro Pro 325 330 Glu Glu Leu Ile Val Phe Cys Gly Leu Leu Tyr Leu Ile Tyr Asp Pro 340 345 350 Ile Lys Lys Phe Gly Asp Glu Asn Thr Ser Ile Met Arg Gly Cys Ala 355 360 365 Ala Ala Glu Arg Phe Tyr Glu Val Leu Asn His Pro Asp Leu His Ser 370 380 Gln Lys Glu Arg Glu Ile Glu Phe Leu Gly Leu Ser Asn Thr Ile Thr Phe Glu Asn Val Ser Phe Gly Tyr Gln Glu Asp Lys His Ile Leu Lys 405 410 415 Asn Leu Ser Phe Thr Leu His Lys Gly Glu Ala Leu Gly Ile Val Gly
420 425 430 Pro Thr Gly Ser Gly Lys Thr Thr Leu Val Lys Leu Leu Pro Arg Leu 435 440 445 Tyr Glu Val Ser Gln Gly Lys Ile Leu Ile Asp Ser Leu Pro Ile Thr Glu Tyr Asn Lys Gly Ser Leu Arg Asn His Ile Ala Cys Val Leu Gln 470 475 Asn Pro Phe Leu Phe Tyr Asp Thr Val Trp Asn Asn Leu Thr Cys Gly
485 490 495 Lys Asp Met Glu Glu Glu Ala Val Leu Glu Ala Leu Lys Arg Ala Tyr 500 505 510 Ala Asp Glu Phe Ile Leu Lys Leu Pro Lys Gly Val His Ser Val Leu 515 520 525 Glu Glu Ser Gly Lys Asn Leu Ser Gly Gly Gln Gln Gln Arg Leu Ala 535 540 Ile Ala Arg Ala Leu Leu Lys Asn Ala Ser Ile Leu Ile Leu Asp Glu 550 555 Ala Thr Ser Ala Leu Asp Ala Ile Ser Glu Asn Tyr Ile Lys Asn Ile 565 570 575 Ile Gly Glu Leu Lys Gly Gln Cys Thr Gln Ile Ile Ile Ala His Lys 585 580 590 Leu Thr Thr Leu Glu His Val Asp Arg Val Leu Tyr Ile Glu Asn Gly 600 Gln Lys Ile Ala Glu Gly Thr Lys Glu Glu Leu Leu Gln Thr Cys Pro 610 620 \ Glu Phe Leu Lys Met Trp Glu Leu Ser Gly Thr Lys Glu Tyr Asn Arg 625 630 635 640 Val Phe Val Pro Asp His Lys Leu Val Ala Asn Pro Thr Asp Met Ala 650 645 Ile Thr Thr

<210> 498 <211> 411 <212> PRT

<213> Chlamydia pneumoniae

<400> 498

Met Ile Pro Thr Met Leu Met Phe Phe Ile Ile Cys Phe Thr Leu Cys 5 10 Ser Gly Phe Ile Ser Leu Ser Gln Ile Ala Leu Phe Ser Leu Pro Thr 20 25 . Ser Leu Ile Ser His Tyr Lys Arg Ser Lys Ser Lys Lys Gln Gln Arg 35 40 45 Val Ala Thr Leu Leu Hes Pro His His Leu Leu Ile Thr Leu Ile 55 Phe Cys Asp Ile Gly Leu Asn Ile Ala Ile Gln Asn Cys Phe Ala Ile 70 75 Leu Phe Gly Asp Ala Ala Ser Trp Trp Phe Thr Val Gly Leu Pro Leu 90 Ala Ile Thr Leu Ile Leu Gly Glu Ile Leu Pro Lys Ala Val Ala Leu 100 105 Pro Phe Asn Thr Gln Ile Ala Ser Ser Val Ala Pro Leu Ile Leu Cys 115 120 125 Val Thr Lys Ile Phe Lys Pro Leu Leu His Trp Gly Ile Val Gly Ile 130 135 140 Asn Tyr Val Val Gln Trp Ile Leu Ser Lys Gln Gln Ile Asp Ile Ile 145 150 150 160 Gln Pro Gln Glu Leu Lys Glu Val Leu Gln Ser Cys Lys Asp Phe Gly 165 170 175 Val Val Asn Gln Glu Glu Ser Arg Leu Leu Tyr Gly Tyr Leu Ser Leu 185 Ser Asp Cys Ser Val Lys Glu Arg Met Gln Pro Arg Gln Asp Ile Leu 200 205 195 Phe Tyr Asp Ile Gln Thr Pro Leu Glu Asn Leu Tyr Leu Leu Phe Ser 210 215 220

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Lys Gln His Cys Ser Arg Val Pro Ile Cys Asn Asp Asn Leu Gln Asn
                   230
                                            235
Leu Leu Gly Ile Cys Thr Ala Arg Ser Leu Leu Leu His Asp Lys Pro 245 \hspace{1cm} 250 \hspace{1cm} 255 \hspace{1cm}
Leu Gln Ser Ser Asp Asp Leu Leu Pro Leu Leu Lys Lys Pro Tyr Tyr 260 265 270
Met Pro Glu Thr Ile Ser Ala Lys Met Ala Leu Cys Gln Met Ala Ala
275 280 285
Glu Asp Glu Thr Leu Gly Met Ile Ile Asp Glu Tyr Gly Ser Ile Glu
290 295 300
Gly Leu Ile Thr Gln Glu Asp Leu Phe Glu Ile Val Ala Gly Glu Ile
                   310 315
Val Asp Gln Arg Asp Asn Lys Ile Leu Tyr Thr Thr Ser Gly Ala Asp 325 330 335
Val Ile Ile Ala Ser Gly Thr Leu Glu Leu Arg Glu Phe Ser Glu Ile 340 345 350
Phe Asp Ile Asn Leu Pro Thr Asn Asn Asn Ile Ala Thr Ile Gly Gly 355 360 365
Trp Leu Ile Glu Gln Ile Gly Thr Ile Pro Thr Thr Gly Met Lys Leu 370 375 380
                      375
Ser Trp Asn Asn Leu Leu Phe Gln Val Leu Asp Ala Ala Pro Asn Arg
385 390 395 400
Ile Arg Arg Val Tyr Ile Arg Lys Leu Tyr Asp
               405
```

<210> 499 <211> 404

<212> PRT

<213> Chlamydia pneumoniae

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Asn Val Ile Gly Ile Ala His Pro Lys Asp Phe Val Asn Lys Ala Leu 245 250 255Asp Glu Pro Leu Ile Asn Asn Leu His Ser Pro Trp Phe Ile Thr Ala 260 265 270 Lys Ser Lys Leu Ile Arg Ile Leu Lys Glu Phe Arg Asp Asn Arg Ser 275 280 285 Ser Val Ala Val Val Leu Asn Ala Ser Gly Glu Pro Ile Gly Ile Leu 295 300 Ser Leu Asn Ala Ile Phe Lys Ile Leu Phe Asn Thr Thr Asn Ile Ala 310 315 His Leu Lys Pro Lys Thr Ile Ser Val Ile Glu Arg Thr Phe Pro Gly 325 330 335 Asn Ser Arg Ile Lys Asp Leu Gln Lys Glu Leu Asp Ile Gln Phe Pro 340 345 Gln Tyr Pro Val Glu Thr Leu Ala Gln Leu Val Leu Gln Leu Leu Asp 355 360 365 355 Ser Pro Ala Glu Val Gly Thr Ser Val Ile Ile Asn Asn Leu Leu Leu 370 375 380 Glu Val Lys Glu Met Ser Leu Ser Gly Ile Lys Thr Val Ser Ile Lys 390 395 Asn Leu Leu Ser

<210> 500 <211> 543

<212> PRT <213> Chlamydia pneumoniae

<400> 500 Met Phe Gly Ser Glu Ser Leu Arg Tyr Gln Leu Leu Ile Gln Asp Phe 10 Ala Lys Val Ser Glu Glu Gly Ile Gly Leu Leu Glu Ser Lys Glu Tyr 20 25 30 Ser Leu Leu Gln Ala Lys Leu Val Leu Arg Ala Leu Ala Gln Asn Ser 35 40 45 Ser Phe Asp Asp Trp Phe Arg Ser Phe Lys Lys Cys Gln Ile Ser Tyr 55 60 Pro Glu Leu Ala His Asp Arg Asp Val Leu Glu Glu Phe Gly Ile Gln 65 70 75 80 Val Leu Arg Glu Gly Ile Glu Asn Pro Ser Val Thr Val Arg Ala Val 85 90 95 90 Ser Val Leu Ala Ile Gly Leu Ala Arg Asp Phe Arg Leu Val Pro Leu 100 105 Phe Arg Leu Val Pro Leu 100Leu Leu Gln Ser Cys Asn Asp Asp Ser Ala Ile Val Arg Ser Leu Ala 115 120 125Leu Gln Val Ala Val Asn Tyr Gly Ser Glu Ser Leu Lys Lys Ala Ile 130 135 140 140 Val Glu Leu Ala Arg Asn Asp Asp Ser Ile His Val Arg Ile Thr Ala 145 150 155 160 Tyr Gln Val Val Ala Leu Leu Gln Ile Glu Glu Leu Leu Pro Phe Leu 165 170 175 Arg Glu Arg Ala Glu Asn Lys Leu Val Asp Ser Val Glu Arg Arg Glu 180 185 190Ala Trp Lys Ala Cys Leu Glu Leu Ser Ser Gln Phe Leu Glu Thr Gly 195 200 205 Val Ala Lys Asp Asp Ile Asp Gln Ala Leu Phe Thr Cys Glu Val Leu 210 215 220 Arg Asn Gly Met Leu Pro Glu Thr Thr Glu Ile Phe Thr Glu Leu Leu 235 240 230 Ser Val Glu His Pro Glu Val Gln Glu Ser Leu Leu Leu Ser Ala Leu

Ala Trp Ser His Gln Leu Gln Asn His Lys Glu Phe Leu Ser Lys Val 260 265 270Arg His Val Met Cys Thr Ser Pro Phe Ala Lys Val Arg Phe Gln Ala 275 280 285 Ala Ala Leu Leu His Leu His Gly Asp Pro Leu Gly Arg Asp Ser Leu 290 295 300 Val Glu Gly Leu Arg Ser Pro Gln Pro Leu Val Cys Glu Ala Ala Ser 305 310 315 320 Ala Ala Leu Cys Ser Leu Gly Ile His Gly Val Pro Leu Ala Lys Glu 325 330 335 His Leu Glu Ser Leu Ser Ser Arg Lys Ala Ala Ala Asa Leu Ser Ile $340 \hspace{1cm} 345 \hspace{1cm} 350$ Leu Leu Val Ser Arg Glu Asp Ile Glu Arg Ala Gly Asp Val Ile Ala Arg Tyr Leu Ser Asn Pro Glu Met Cys Trp Ala Ile Glu Tyr Phe 370 380 Leu Trp Asp Ala Gln Trp Asn Leu Arg Gly Asp Thr Phe Pro Leu Tyr 385 390 395 400 Ser Asp Met Ile Lys Arg Glu Ile Gly Arg Lys Leu Ile Arg Leu Leu 405 415 Ala Val Ala Arg Tyr Ser Gln Ala Lys Ala Val Thr Ala Thr Phe Leu 420 425 430Ser Gly Gln Gln Ala Gln Gly Trp Ser Phe Phe Ser Gly Met Phe Trp 435 440 445Glu Glu Gly Asp Val Lys Thr Ser Glu Asp Leu Val Thr Asp Ala Cys
450 460 Phe Ala Ala Lys Leu Glu Gly Ala Leu Ala Ser Leu Cys Gln Lys Lys 465 470 475 480 Asp Gln Ala Ser Leu Gln Arg Val Ser Gln Leu Tyr Asn Asp Ser Arg 485 490 495 Trp Gln Asp Lys Leu Ala Ile Leu Glu Ser Val Ala Phe Ser Glu Asn $500 \hspace{1.5cm} 505 \hspace{1.5cm} 510$ Leu Asp Ala Val Pro Phe Leu Leu Asp Cys Cys His His Glu Ala Pro 515 520 525 Ser Leu Arg Ser Ala Ala Ala Gly Ala Leu Phe Ser Ile Phe Lys

<400> 501

Met Ser Phe Lys Arg Phe Leu Gln Gln Ile Pro Val Arg Ile Cys Leu Leu Ile Ile Tyr Leu Tyr Gln Trp Leu Ile Ser Pro Leu Leu Gly Ser 25 Cys Cys Arg Phe Phe Pro Ser Cys Ser His Tyr Ala Glu Gln Ala Leu 35 40 45 Lys Ser His Gly Phe Leu Met Gly Cys Trp Leu Ser Ile Lys Arg Ile 50 55 60 Gly Lys Cys Gly Pro Trp His Pro Gly Gly Ile Asp Met Val Pro Lys 65 70 75 80Thr Ala Leu Glu Val Leu Glu Pro Tyr Gln Glu Ile Asp Gly Gly 85 90 Asp Ser Ser His Phe Ser Glu 100

<210> 501

<211> 103 <212> PRT

<213> Chlamydia pneumoniae

<210> 502

<211> 362 <212> PRT

<213> Chlamydia pneumoniae

<400> 502 Met Ala Phe Lys Arg Lys Thr Arg Trp Leu Trp Gln Val Leu Ile Leu 10 Ser Val Gly Leu Asn Met Leu Phe Leu Leu Leu Phe Tyr Ser Ala Ile 25 30 Phe Arg Lys Asp Ile Tyr Lys Leu His Leu Phe Ser Gly Pro Leu Ile 40 4.5 Ala Lys Ser Ser Arg Lys Val Tyr Leu Ser Glu Asp Phe Leu Asn Glu 55 60 Ile Ser Gln Ala Ser Leu Asp Asp Leu Ile Ser Leu Phe Lys Asp Glu 65 70 75 80 Arg Tyr Met Tyr Gly Arg Pro Ile Lys Leu Trp Ala Leu Ser Val Ala 85 90 Ile Ala Ser His His Ile Asp Ile Thr Pro Val Leu Ser Lys Pro Leu 105 110 Thr Tyr Thr Glu Leu Lys Gly Ser Ser Val Arg Trp Leu Leu Pro Asn 115 120 125Ile Asp Leu Lys Asp Phe Pro Val Ile Leu Asp Tyr Leu Arg Cys His 135 140 Lys Tyr Pro Tyr Thr Ser Lys Gly Leu Phe Leu Leu Ile Glu Lys Met 150 155 Val Glu Glu Gly Trp Val Asp Glu Asp Cys Leu Tyr His Phe Cys Ser 165 170 175 Thr Pro Glu Phe Leu Tyr Leu Arg Thr Leu Leu Val Gly Ala Asp Val 180 185 190 Gln Ala Ser Ser Val Ala Ser Leu Ala Arg Met Val Ile Arg Cys Gly 200 Ser Glu Arg Phe Phe His Phe Cys Asn Glu Glu Ser Arg Thr Ser Met 215 220 Ile Ser Ala Thr Gln Arg Gln Lys Val Leu Lys Ser Tyr Leu Asp Cys 230 235 Glu Glu Ser Leu Ala Ala Leu Leu Leu Leu Val His Asp Ser Asp Val 245 250 Val Leu His Glu Phe Cys Asp Glu Asp Leu Glu Lys Val Ile Arg Leu 260 265 Met Pro Gln Glu Ser Pro Tyr Ser Gln Asn Phe Phe Ser Arg Leu Gln 275 280 285 His Ser Pro Arg Arg Glu Leu Ala Cys Met Ser Thr Gln Arg Val Glu 295 300 Ala Pro Arg Val Gln Glu Asp Gln Asp Glu Glu Tyr Val Val Gln Asp 310 315 Gly Asp Ser Leu Trp Leu Ile Ala Lys Arg Phe Gly Ile Pro Met Asp 325 330 335 Lys Ile Ile Gln Lys Asn Gly Leu Asn His His Arg Leu Phe Pro Gly 340 345 Lys Val Leu Lys Leu Pro Ala Lys Gln Ser

<210> 503

355

<211> 582

<212> PRT

<213> Chlamydia pneumoniae

<400> 503

Met Ser Gly Lys Lys Asp Gly Val Arg Gly Met Ile Phe Val Pro Leu 15 10 15 15 Ser Ile Leu Val Leu Ile Phe Leu Pro Leu Pro Gln Ile Leu Leu Asp 20 25 30 Phe Gly Leu Cys Ile Ser Phe Ala Leu Ser Leu Leu Thr Val Cys Trp

		35					40					45			
Val	Phe 50	Thr	Leu	Asn	Ser	Ser 55	Asn	Ser	Ala	ГÀЗ	Leu 60	Phe	Pro	Pro	Phe
Phe 65	Leu	Tyr	Leu	Cys	Leu 70	Leu	Arg	Leu	Gly	Leu 75	Asn	Leu	Ala	Ser	Thr 80
Arg	Trp	Ile	Val	Ser 85	Ser	Gly	Thr	Ala	Ser 90	Ser	Leu	Ile	Val	Ser 95	Leu
Gly	Ser	Phe	Phe 100	Ser	Leu	Gly	Ser	Leu 105	Trp	Ala	Ala	Thr	Phe 110	Ala	Cys
Leu	Leu	Leu 115	Phe	Phe	Val	Asn	Phe 120	Leu	Met	Val	Ser	Lys 125	Gly	Ser	Glu
Arg	Ile 130	Ala	Glu	Val	Arg	Ser 135	Arg	Phe	Phe	Leu	Glu 140	Ala	Leu	Pro	Ala
Lys 145	Gln	Met	Ala	Leu	Asp 150	Ser	Asp	Leu	Val	Ser 155	Gly	Arg	Ala	Ser	Tyr 160
			Lys	Lys 165	Gln	Lys	Asn	Ala	Leu 170	Ile	Glu	Glu	Gly	Asp 175	Phe
Phe	Ser	Ala	Met 180	Glu	Gly	Val	Phe	Arg 185	Phe	Val	Lys	Gly	Asp 190	Ala	Ile
Ile	Ser	Cys 195	Ile	Leu	Leu	Leu	Val 200	Asn	Val			205		-	
Tyr	210		Ser		-	215					220		Thr		
225	_		Leu		230					235					240
Ala			Leu	245					250					255	Asn
Tyr			Glu 260	-	-	-		265	-				270		Val
Ser		275	Ile				280	-				285		-	
Pro	290		Leu			295			-		300	-	_	-	
305	Pro				310				Glu	315				Tyr	320
	_		Cys	325	_				330			-		335	-
Arg			Ser 340					345	-		_		35Ō		
Val		355	Ser				360					365			
-	Gln 370			-		375					380		Val		
385		-	Asn		390					395					400
	_		Glu	405					410	-				415	-
Ile			Lys 420	-				425					430		-
		435	Arg				440		-			445	-		
	450		Ala			455					460				
465			Val		470					475		_	_		480
			Lys	485					11e 490			_	Phe	495	Val
Glu			11e 500					505	-				510		
GIU	Asn	Val 515	Ile	Arg	Arg	val	520	ser	Leu	Leu	GIu	Arg 525	Ser	Va⊥	rne

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Lys Asp Phe Arg Ala Ile Val Thr Ser Cys Glu Thr Arg Phe Glu Met
                       535
Lys Lys Met Leu Asp Pro His Phe Pro Asp Leu Leu Val Leu Ser His
                            555
                   550
Asp Glu Leu Pro Lys Glu Ile Pro Ile Ser Phe Leu Gly Ile Val Ser
              565
Asp Glu Val Leu Val Pro
           580
<210> 504
<211> 435
<212> PRT
<213> Chlamydia pneumoniae
<400> 504
Met Phe Ser Arg Trp Ile Thr Leu Phe Leu Leu Phe Ile Ser Leu Thr
                                   10
Gly Cys Ser Ser Tyr Ser Ser Lys His Lys Gln Ser Leu Ile Ile Pro
       20
                               25
Ile His Asp Asp Pro Val Ala Phe Ser Pro Glu Gln Ala Lys Arg Ala
                          40
Met Asp Leu Ser Ile Ala Gln Leu Leu Phe Asp Gly Leu Thr Arg Glu
                      55
Thr His Arg Glu Ser Asn Asp Leu Glu Leu Ala Ile Ala Ser Arg Tyr
65 70 75 80
Thr Val Ser Glu Asp Phe Cys Ser Tyr Thr Phe Phe Ile Lys Asp Ser
               85
Ala Leu Trp Ser Asp Gly Thr Pro Ile Thr Ser Glu Asp Ile Arg Asn
                            105
Ala Trp Glu Tyr Ala Gln Glu Asn Ser Pro His Ile Gln Ile Phe Gln
                                              125
      115
                          120
Gly Leu Asn Phe Ser Thr Pro Ser Ser Asn Ala Ile Thr Ile His Leu
   130
                       135
                                          140
Asp Ser Pro Asn Pro Asp Phe Pro Lys Leu Leu Ala Phe Pro Ala Phe
                 150
                                    155
Ala Ile Phe Lys Pro Glu Asn Pro Lys Leu Phe Ser Gly Pro Tyr Thr
              165
                                  170
Leu Val Glu Tyr Phe Pro Gly His Asn Ile His Leu Lys Lys Asn Pro
                                                 190
          180
                              185
Asn Tyr Tyr Asp Tyr His Cys Val Ser Ile Asn Ser Ile Lys Leu Leu
                          200
                                             205
Ile Ile Pro Asp Ile Tyr Thr Ala Ile His Leu Leu Asn Arg Gly Lys
                      215
                                         220
Val Asp Trp Val Gly Gln Pro Trp His Gln Gly Ile Pro Trp Glu Leu
225 230 235 240
His Lys Gln Ser Gln Tyr His Tyr Tyr Thr Tyr Pro Val Glu Gly Ala
245 250 255
Phe Trp Leu Cys Leu Asn Thr Lys Ser Pro His Leu Asn Asp Leu Gln 260 265 270
Asn Arg His Arg Leu Ala Thr Cys Ile Asp Lys Arg Ser Ile Ile Glu
275 280 285
Glu Ala Leu Gln Gly Thr Gln Gln Pro Ala Glu Thr Leu Ser Arg Gly
Ala Pro Gln Pro Asn Gln Tyr Lys Lys Gln Lys Pro Leu Thr Pro Gln 305 310 315 320
Glu Lys Leu Val Leu Thr Tyr Pro Ser Asp Ile Leu Arg Cys Gln Arg
               325
                                330
Ile Ala Glu Ile Leu Lys Glu Gln Trp Lys Ala Ala Gly Ile Asp Leu 340 345
                             345
Ile Leu Glu Gly Leu Glu Tyr His Leu Phe Val Asn Lys Arg Lys Val
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PCT/US01/23121 301

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Gln Asp Tyr Ala Ile Ala Thr Gln Thr Gly Val Ala Tyr Tyr Pro Gly
    370
                         375
Ala Asn Leu Ile Ser Glu Glu Asp Lys Leu Leu Gln Asn Phe Glu Ile
                    390
                                         395
Ile Pro Ile Tyr Tyr Leu Ser Tyr Asp Tyr Leu Thr Gln Asp Phe Ile 405 410 415
Glu Gly Val Ile Tyr Asn Ala Ser Gly Ala Val Asp Leu Lys Tyr Thr 420 \hspace{1.5cm} 425 \hspace{1.5cm} 430
Tyr Phe Pro
       435
<210> 505
<211> 171
<212> PRT
<213> Chlamydia pneumoniae
<400> 505
Met Lys Lys Leu Leu Phe Ser Thr Phe Leu Leu Val Leu Gly Ser Thr
Ser Ala Ala His Ala Asn Leu Gly Tyr Val Asn Leu Lys Arg Cys Leu
                                 25
Glu Glu Ser Asp Leu Gly Lys Lys Glu Thr Glu Glu Leu Glu Ala Met
                            40
Lys Gln Gln Phe Val Lys Asn Ala Glu Lys Ile Glu Glu Glu Leu Thr
                       55
Ser Ile Tyr Asn Lys Leu Gln Asp Glu Asp Tyr Met Glu Ser Leu Ser
65 70 75 80
Asp Ser Ala Ser Glu Glu Leu Arg Lys Lys Phe Glu Asp Leu Ser Gly
Glu Tyr Asn Ala Tyr Gln Ser Gln Tyr Tyr Gln Ser Ile Asn Gln Ser 100 105 110
Asn Val Lys Arg Ile Gln Lys Leu Ile Gln Glu Val Lys Tle Ala Ala
115 120 125
Glu Ser Val Arg Ser Lys Glu Lys Leu Glu Ala Ile Leu Asn Glu Glu
130 135 140
Ala Val Leu Ala Ile Ala Pro Gly Thr Asp Lys Thr Thr Glu Ile Ile
145 150 150 160
Ala Ile Leu Asn Glu Ser Phe Lys Lys Gln Asn
                165
<210> 506
<211> 360
<212> PRT
<213> Chlamydia pneumoniae
<400> 506
Met Ser Glu Ala Pro Val Tyr Thr Leu Lys Gln Leu Ala Glu Leu Leu
                                     10
Gln Val Glu Val Gln Gly Asn Ile Glu Thr Pro Ile Ser Gly Val Glu
                                25
Asp Ile Ser Gln Ala Gln Pro His His Ile Ala Phe Leu Asp Asn Glu
                         40
Lys Tyr Ser Ser Phe Leu Lys Asn Thr Lys Ala Gly Ala Ile Ile Leu
                     55
                                             60
Ser Arg Ser Gln Ala Met Gln His Ala His Leu Lys Lys Asn Phe Leu
                    70
Ile Thr Asn Glu Ser Pro Ser Leu Thr Phe Gln Lys Cys Ile Glu Leu
                                   90
              85
Phe Ile Glu Pro Val Thr Ser Gly Phe Pro Gly Ile His Pro Thr Ala
                                105
Val Ile His Pro Thr Ala Arg Ile Glu Lys Asn Val Thr Ile Glu Pro
```

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120
Tyr Val Val Ile Ser Gln His Ala His Ile Gly Ser Asp Thr Tyr Ile 130 $140\,
Gly Ala Gly Ser Val Ile Gly Ala His Ser Val Leu Gly Ala Asn Cys
145 150 155 160
Leu Ile His Pro Lys Val Val Ile Arg Glu Arg Val Leu Met Gly Asn
165 170 175
Arg Val Val Val Gln Pro Gly Ala Val Leu Gly Ser Cys Gly Phe Gly 180 185 190
Tyr Ile Thr Asn Ala Phe Gly His His Lys Pro Leu Lys His Leu Gly
195 200 205
Tyr Val Ile Val Gly Asp Asp Val Glu Ile Gly Ala Asn Thr Thr Ile
                     215
Asp Arg Gly Arg Phe Lys Asn Thr Val Ile His Glu Gly Thr Lys Ile
225 230 235 240
Ile Ile Val Ala Gln Ala Gly Ile Ala Gly Ser Thr Lys Ile Gly Glu 260 265 270
His Val Ile Ile Gly Gly Gln Thr Gly Ile Thr Gly His Ile Ser Ile
275 280 285
Ala Asp His Val Ile Met Ile Ala Gln Thr Gly Val Thr Lys Ser Ile
290 295 300
Thr Ser Pro Gly Ile Tyr Gly Gly Ala Pro Ala Arg Pro Tyr Gln Glu
305 310 315 320
Thr His Arg Leu Ile Ala Lys Ile Arg Asn Leu Pro Lys Thr Glu Glu
325 330 335
Arg Leu Ser Lys Leu Glu Lys Gln Val Arg Asp Leu Ser Thr Pro Ser
         340 345
Leu Ala Glu Ile Pro Ser Glu Ile
       355 , 360
<210> 507
<211> 399
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<212> PRT <213> Chlamydia pneumoniae

<400> 507

Met Ala Ala Ser Gly Gly Thr Gly Gly Leu Gly Gly Thr Gln Gly Val Asn Leu Ala Ala Val Glu Ala Ala Ala Ala Lys Ala Asp Ala Ala Glu 25 Val Val Ala Ser Gln Glu Gly Ser Glu Met Asn Met Ile Gln Gln Ser 35 40 45 Gln Asp Leu Thr Asn Pro Ala Ala Ala Thr Arg Thr Lys Lys Lys Glu 50 55 60 Glu Lys Phe Gln Thr Leu Glu Ser Arg Lys Lys Gly Glu Ala Gly Lys 65 70 75 80 Ala Glu Lys Lys Ser Glu Ser Thr Glu Glu Lys Pro Asp Thr Asp Leu 85 90 95 Ala Asp Lys Tyr Ala Ser Gly Asn Ser Glu Ile Ser Gly Gln Glu Leu 100 105 110 Arg Gly Leu Arg Asp Ala Ile Gly Asp Asp Ala Ser Pro Glu Asp Ile 115 120 125 Leu Ala Leu Val Gln Glu Lys Ile Lys Asp Pro Ala Leu Gln Ser Thr 130 135 140 Ala Leu Asp Tyr Leu Val Gln Thr Thr Pro Pro Ser Gln Gly Lys Leu 145 150 155 160 Lys Glu Ala Leu Ile Gln Ala Arg Asn Thr His Thr Glu Gln Phe Gly 165 170 175 Arg Thr Ala Ile Gly Ala Lys Asn Ile Leu Phe Ala Ser Gln Glu Tyr

180 185 Ala Asp Gln Leu Asn Val Ser Pro Ser Gly Leu Arg Ser Leu Tyr Leu 195 200 205 Glu Val Thr Gly Asp Thr His Thr Cys Asp Gln Leu Leu Ser Met Leu 215 220 Gln Asp Arg Tyr Thr Tyr Gln Asp Met Ala Ile Val Ser Ser Phe Leu 225 230 235 240 Met Lys Gly Met Ala Thr Glu Leu Lys Arg Gln Gly Pro Tyr Val Pro 245 250 255 Ser Ala Gln Leu Gln Val Leu Met Thr Glu Thr Arg Asn Leu Gln Ala 260 265 270 Val Leu Thr Ser Tyr Asp Tyr Phe Glu Ser Arg Val Pro Ile Leu Leu 280 285 Asp Ser Leu Lys Ala Glu Gly Ile Gln Thr Pro Ser Asp Leu Asn Phe 295 300 Val Lys Val Ala Glu Ser Tyr His Lys Ile Ile Asn Asp Lys Phe Pro 310 315 Thr Ala Ser Lys Val Glu Arg Glu Val Arg Asn Leu Ile Gly Asp Asp 325 330 335 Val Asp Ser Val Thr Gly Val Leu Asn Leu Phe Phe Ser Ala Leu Arg 340 345 350Gln Thr Ser Ser Arg Leu Phe Ser Ser Ala Asp Lys Arg Gln Gln Leu 355 360 365 Gly Ala Met Ile Ala Asn Ala Leu Asp Ala Val Asn Ile Asn Asn Glu 375 380 Asp Tyr Pro Lys Ala Ser Asp Phe Pro Lys Pro Tyr Pro Trp Ser <210> 508 <211> 224 <212> PRT <213> Chlamydia pneumoniae <400> 508 Met Thr Ser Trp Ile Glu Leu Leu Asp Lys Gln Ile Glu Asp Gln His 1 5 10 15 Met Leu Lys His Glu Phe Tyr Gln Arg Trp Ser Glu Gly Lys Leu Glu Lys Gln Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr Leu His Ile Lys 40 Ala Phe Pro Cys Tyr Leu Ser Ala Leu His Ala Arg Cys Asp Asp Leu 55 Gln Ile Arg Arg Gln Ile Leu Glu Asn Leu Met Asp Glu Glu Ala Gly 70 7.5 Asn Pro Asn His Ile Asp Leu Trp Arg Gln Phe Ala Leu Ser Leu Gly 85 90 Val Ser Glu Glu Glu Leu Ala Asn His Glu Phe Ser Gln Ala Ala Gln 100 105 110 Asp Met Val Ala Thr Phe Arg Arg Leu Cys Asp Met Pro Gln Leu Ala Val Gly Leu Gly Ala Leu Tyr Thr Tyr Glu Ile Gln Ile Pro Gln Val 130 135 140 Cys Val Glu Lys Ile Arg Gly Leu Lys Glu Tyr Phe Gly Val Ser Ala 145 150 155 160Arg Gly Tyr Ala Tyr Phe Thr Val His Gln Glu Ala Asp Ile Lys His 170 Ala Ser Glu Glu Lys Glu Met Leu Gln Thr Leu Val Gly Arg Glu Asn 180 185 190 Pro Asp Ala Val Leu Gln Gly Ser Gln Glu Val Leu Asp Thr Leu Trp

Asn Phe Leu Ser Ser Phe Ile Asn Ser Thr Glu Pro Cys Ser Cys Lys

WO 02/08267 PCT/US01/23121

220

304

215

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<211> 353 <212> PRT <213> Chlamydia pneumoniae

<400> 510

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 Leu Ile Leu Val Gly Ile Phe Ala Arg Ala Fro Arg Gly Asp Thr

 25
 30

 26
 25

 27
 Asp Glu Ala Ile Ile Tyr Ser Asn Gln

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 Cys Asn Glu Asp Met Arg Lys Ile Leu Cys Asp Ala Ile Glu His Ala

 50
 55

 Asp Glu Glu Ile Phe Lu Arg Ile Tyr Asn Leu Ser Glu Pro Lys Ile

 65
 70

 61n Gln Ser Leu Thr Arg Gln Ala Gln Ala Lys Asn Lys Val Thr Ile

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 67
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 7

Val Thr Leu Val Glu Gln Pro Pro Ala Gly Arg Lys Leu Met His Gln 115 120 125 Lys Ala Leu Ser Ile Asp Lys Lys Asp Ala Trp Leu Gly Ser Ala Asn 130 140 Tyr Thr Asn Leu Ser Leu Arg Leu Asp Asn Asn Leu Ile Leu Gly Met 150 155 His Ser Ser Glu Leu Cys Asp Leu Ile Ile Thr Asn Thr Ser Gly Asp 165 170 175 Phe Ser Ile Lys Asp Gln Thr Gly Lys Tyr Phe Val Leu Pro Gln Asp 180 185 190 Arg Lys Ile Ala Ile Gln Ala Val Leu Glu Lys Ile Gln Thr Ala Gln 195 200 205 Lys Thr Ile Gln Val Ala Met Phe Ala Leu Thr His Ser Glu Ile Ile 210 215 220 Gln Ala Leu His Gln Ala Lys Gln Arg Gly Ile His Val Asp Ile Ile 225 230 235 240 Ile Asp Arg Ser His Ser Lys Leu Thr Phe Lys Gln Leu Arg Gln Leu 245 250 255 Asn Ile Asn Lys Asp Phe Val Ser Ile Asn Thr Ala Pro Cys Thr Leu 260 265 270His His Lys Phe Ala Val Ile Asp Asn Lys Thr Leu Leu Ala Gly Ser 275 280 285 Ile Asn Trp Ser Lys Gly Arg Phe Ser Leu Asn Asp Glu Ser Leu Ile 290 295 300 Ile Leu Glu Asn Leu Thr Lys Gln Gln Asn Gln Lys Leu Arg Met Ile 310 315 Trp Lys Asp Leu Ala Lys His Ser Glu His Pro Thr Val Asp Asp Glu 325 330 335 Glu Lys Glu Ile Ile Glu Lys Ser Leu Pro Val Glu Glu Glu Glu Ala 340 345

<210> 511 <211> 186 <212> PRT

<213> Chlamydia pneumoniae

<400> 511

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Glu Lys Ser His Ala Gln Asp Glu Asn Gln
           180
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<220>
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Ala Asp Asn Ile Arg Leu Gly Gln Met Thr Thr Val Leu Lys Lys Asp
                                25
Glu Val Ile Ile Gly Thr Asp Thr Thr Pro Thr Val Thr Lys Phe Ser
       35
                           40
Gly Asp Lys Gly Ile Val Ile Thr Thr Asp Ser Thr Ile Thr Pro Ser
                       55
Ser Thr Thr Phe Ser Leu Asp Met Glu Ala Val Ile Lys Glu Val Thr
                   70
                                       75
Asp Lys Ile Leu Thr Gln Ile Glu Asp Glu Leu Val Lys Asp Ile Ile
                85
                                    90
Lys Asn Ile Thr Gln Ser Leu Ile Glu Glu Val Ile Lys Lys Ile His
           100
                               105
                                                   110
Ile Asp Pro Ser Phe Ser Tyr Ser Arg Ala Phe Lys Asp Val Asn Ile
115 120 125
Thr Asn Lys Ile Gln Cys Asn Gly Leu Phe Thr Lys Glu Asn Ile Gly
                       135
                                           140
Asn Leu Asp Gly Gly Thr Glu Ile Ala Ser Ser Ser Val Thr Pro Asp
                    150
                                        155
Asn Ala Asn Ser Met Phe Leu Ile Cys Ala Asp Ile Ile Ala Thr Arg
               165
                                  170
Met Glu Gly Thr Val Ala Leu Ala Leu Val Lys Glu Gly Asp Leu Ser
           180
                               185
                                                   190
Pro Cys Ser Ile Ser Tyr Gly Tyr Ser Ala Gly Tyr Pro Asn Ile Ile
       195
                            200
                                               205
Ser Leu Arg Ala Thr Val Gly Asn Lys Thr Thr Ala Pro Val Lys Phe
                        215
                                            220
Ser Leu Arg Ala Gly Gly Met Asp Ser Gly Val Val Trp Val Asn Ala
                    230
Met Pro Asn Gly Glu Lys Ile Leu Gly Val Asp Ala Val Ser Lys Ile
                245
                                   250
Thr Ile Leu Glu Val Lys Pro Gln Thr Asn Gly Thr Xaa Xaa Xaa Xaa
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Phe Xaa Xaa Xaa
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Met Val Glu Val Glu Glu Lys His Tyr Thr Ile Val Lys Arq Asn Gly
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Met Phe Val Pro Phe Asn Gln Asp Arg Ile Phe Gln Ala Leu Glu Ala
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Ala	Phe	Arg 35	Asp	Thr	Arg	Ser	Leu 40	Glu	Thr	Ser	Ser	Pro 45	Leu	Pro	Lys
Asp	Leu 50	Glu	Glu	Ser	Ile	Ala 55	Gln	Ile	Thr	His	Lys 60	Val	Val	Lys	Glu
Val 65	Leu	Ala	Lys	Ile	Ser 70	Glu	Gly	Gln	Val	Val 75	Thr	Val	Glu	Arg	Ile 80
Gln	qzA	Leu	Val	85				-	90		-			95	
	Arg	-	100					105	Gln				110	-	_
	Ser	115					120					125			
-	Phe 130				-	135					140	-			
145	Thr				150				Pro	155					160
	Asn	-		165		-			170	-				175	
	Glu		180					185					190		_
	Leu	195					200					205			
	Arg 210 Glu					215					220			-	
225	Gly		Thr		230				Thr	235				-	240
Ser	-		Cys	245			_	-	250	-			-	255	
	Asp		260					265					270		Glu
Val		275	Ala				280		Arg			285			Glu
	290 Asp					295			_		300			_	
305	-	-			310					315					320
	Glu			325	_				330					335	
His	-	-	His 340		-		-	345			_		350	-	
	Asn	355					360					365			
	370					375			Ser	-	380	_			
385	Tyr				390				Glu	395					400
	Gln			405		-			410	-				415	
	Glu		420					425			-		430		
	Phe	435	_				440					445		_	
	His 450					455					460				
465	Ser				4.70				_	475					480
•	Ala		Gly	485	_			_	490	-		_		495	
Ala	Val		Lys 500					505					510		
Ile	Lys	Val	Ala	Asn	Asp	Thr	Ala	Ile	Ala	Val	Asn	Gln	Gly	Gly	Lys

_	_	515			_		520					525			
	Lys 530	_				535					540				
545	Asp				550					555					560
Thr	His	Asp	Ile	Asn 565	Thr	Ala	Ser	Trp	Ile 570	Pro	Asp	Leu	Phe	Phe 575	Lys
Arg	Leu	Glu	Lys 580	Lys	Gly	Met	Trp	Thr 585	Leu	Phe	Ser	Pro	Asp 590	Asp	Val
Pro	Gly	Leu 595	His	Glu	Ala	Tyr	Gly 600	Leu	Glu	Phe	Glu	Lys 605	Leu	Tyr	Glu
Glu	Tyr 610	Glu	Arg	Lys	Val	Glu 615	Ser	Gly	Glu	Ile	Arg 620	Leu	Tyr	Lys	Lys
Val 625	Glu	Ala	Glu	Val	Leu 630	Trp	Arg	Lys	Met	Leu 635	Ser	Met	Leu	Tyr	Glu 640
Thr	Gly	His	Pro	Trp 645	Ile	Thr	Phe	Lys	Asp 650	Pro	Ser	Asn	Ile	Arg 655	Ser
Asn	Gln	Asp	His 660	Val	Gly	Val	Val	Arg 665	Суз	Ser	Asn	Leu	Cys 670	Thr	Glu
Ile	Leu	Leu 675	Asn	Cys	Ser	Glu	Ser 680	Glu	Thr	Ala	Val	Cys 685	Asn	Leu	Gly
Ser	11e 690	Asn	Leu	Val	Glu	His 695	Ile	Arg	Asn	Asp	Lys 700	Leu	Asp	Glu	Glu
705	Leu	-			710		Ile			715			-		720
Ile	Asp	Leu	Asn	Phe 725	Tyr	Pro	Thr	Pro	Glu 730	Ala	Lys	Gln	Ala	Asn 735	Leu
	His	-	740		-		_	745		_			750		
	Glu	755					760					765			Ser
Asp	Glu 770	Cys	Ser	Glu	Ile	Ile 775	Ala	Tyr	Tyr	Ala	Ile 780	Leu	Ala	Ser	Ser
785	Leu		_		790					795	_		_		800
	Asp		_	805					810					815	
	Arg		820					825					830		_
Trp		835					Tle 840					845			
	Val 850					855					860				_
865	Thr				870				-	875					880
	Leu		_	885					890		-			895	-
	Lys		900					905				_	910		
	Phe	915					920					925			
-	Lys 930					935					940		-		
945	Cys				950					955		-			960
	Leu	-		965			_		970	-				975	Tyr
	Thr		980					985					990		
Gln	Ala	Ala 995	Thr	Ser	Val	Glu	Lys 1000		Phe	Ile	Asp	11e 1005		Lys	Arg

Gly Ile Gln Pro Arg Trp Met Lys Asn Lys Ser Ala Ser Thr Ser Ile 1010 1015 1020 Val Val Glu Arg Lys Thr Thr Pro Val Cys Ser Met Glu Glu Gly Cys 1025 1030 1035 1040 Glu Ser Cvs Gln

<210> 514 <211> 346 <212> PRT <213> Chlamydia pneumoniae

<400> 514

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Ile Asp Tyr Val Arg His Ile Ala Asp Arg Arg Leu Glu Arg Ile Gly 295 300

Leu Lys Pro Ile Tyr His Ser Arg Asn Pro Phe Pro Trp Met Ser Glu 305 310 315 320 Thr Met Asp Leu Asn Lys Glu Lys Asn Phe Phe Glu Thr Arg Val Thr 325 330 335

Glu Tyr Gln Thr Ala Gly Asn Leu Ser Trp 340

<210> 515

<211> 327 <212> PRT <213> Chlamydia pneumoniae

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145 150 150 160 150 155 Lys Phe Pro Glu Trp Ser Ala Glu Phe Lys Ala Asn Ser Glu Glu Leu 165 170 175 Val Cys Glu Met Ser Ile Leu Asp Ser Trp Ala Lys Gln Cys Leu Ser 180 185 190 Thr Ile Pro Glu Asn Leu Arg Tyr Leu Val Ser Gly His Asn Ala Phe 200 205 Ser Tyr Phe Thr Arg Arg Tyr Leu Ala Thr Pro Glu Glu Val Ala Ser 215 210 220 Gly Ala Trp Arg Ser Arg Cys Ile Ser Pro Glu Gly Leu Ser Pro Glu 230 235 Ala Gln Ile Ser Val Arg Asp Ile Met Ala Val Val Asp Tyr Ile Asn 250 245 Glu His Asp Val Ser Val Val Phe Pro Glu Asp Thr Leu Asn Gln Asp 260 270 Ala Leu Lys Lys Ile Val Ser Ser Leu Lys Lys Ser His Leu Val Arg 275 280 285 Leu Ala Gln Lys Pro Leu Tyr Ser Asp Asn Val Asp Asp Asn Tyr Phe 300 295 Ser Thr Phe Lys His Asn Val Cys Leu Ile Thr Glu Glu Leu Gly Gly 310 315 Val Ala Leu Glu Cys Gln Årg

<210> 516 <211> 101

<212> PRT

<213> Chlamydia pneumoniae

325

<400> 516

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Phe Ser Lys Thr Glu
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<211> 261
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<213> Chlamydia pneumoniae
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           20
                               25
Lys Val Leu Leu Val Asp Leu Asp Pro Gln Ala Asn Leu Thr Thr Gly
Leu Gly Val Gln Ser Cys Tyr Glu Ser Asn Leu Asn Asp Ile Phe Arg 50 55 60
Ser Ser Gly Asn Val Arg Asp Ile Ile Gln Asp Thr Lys Ile Glu Asn
                                       75
                 70
Leu His Ile Val Pro Ser Ser Ile Leu Ile Glu Glu Phe Arg Glu Phe
                                  90
             8.5
Asn Arg Asn Ser Val Leu Asp Thr Ser His Leu Arg Ser Ser Leu Gln 100 105 110
Leu Ile Glu Ser Asn Tyr Asp Leu Cys Ile Leu Asp Thr Pro Pro Ser
                            120
                                                125
Leu Gly Thr Leu Thr Glu Glu Ala Phe Ile Ala Ser Asp His Leu Ile
                       135
Val Cys Leu Thr Pro Glu Pro Phe Ser Ile Leu Gly Leu Gln Lys Ile
                   150
                                       155
Lys Glu Phe Cys Ser Val Leu Pro Lys Lys Lys Asp Leu Ser Val Leu
                165
                                    170
Gly Ile Val Phe Ser Phe Trp Asp Gly Arg Asn Ser Thr Asn Ser Thr
                              185
Tyr Leu Asn Ile Ile Glu Ser Ile Tyr Glu Gly Lys Val Leu Ser Ser
195 200 205
Lys Val Arg Arg Asp Ile Thr Leu Ser Arg Ser Leu Leu Lys Glu Thr
Ser Ile Ala Asn Ala Tyr Pro Asn Ser Arg Ala Ser His Asp Ile Leu
225 230 235 240
Arg Leu Thr Lys Glu Ile Glu Asp Lys Leu Phe Asn Lys Glu Met Ser
                245
Ala Gln Glu Val Leu
            260
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<211> 526
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Lys Leu Ile Gly Thr Ser Pro Lys His Gly Ile Tyr Leu Pro Leu Phe
         20
Ser Ile His Thr Lys Asn Ser Cys Gly Ile Gly Glu Phe Leu Asp Leu
                          40
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Ile Pro Leu Ile Ser Trp Cys Gln Lys Gln Gly Phe Ser Val Ile Gln

Leu 65	Leu	Pro	Leu	Asn	Asp 70	Thr	Gly	Glu	Asp	Thr 75	Ser	Pro	Tyr		Ser .80
Ile	Ser	Ser	Val	Ala 85	Leu	Asn	Pro	Leu	Phe 90	Leu	Ser	Leu	Ser	Ser 95	Leu
Pro	Asn	Ile	Asp 100	Thr	Ile	Pro	Glu	Val 105	Ala	Lys	Lys	Leu	Gln 110	Asp	Met
His	Glu	Leu 115	Cys	Ser	Thr	Pro	Ser 120	Val	Ser	Tyr	Thr	Gln 125	Val	Lys	G1u
Lys	Lys 130	Trp	Ala	Phe	Leu	Arg 135	Glu	Tyr	Tyr	Gln	Lys 140	Cys	Cys	Lys	Ser
Ser 145	Leu	Glu	Gly	Asn	Ser 150	Asn	Phe	Ser	Glu	Phe 155	Leu	Glu	Ser	Glu	Arg 160
Tyr	Trp	Leu	Tyr	Pro 165	Tyr	Gly	Thr	Phe	Arg 170	Ala	Ile	Lys	His	His 175	Met
His	Gly	Glu	Pro 180	Ile	Asn	Asn	Trp	Pro 185	Lys	Ser	Leu	Thr	Asp 190	Gln	Glu
Asn	Phe	Pro 195	Asp	Leu	Thr	Ьуз	Lys 200	Phe	His	Asp	Glu	Val 205	Leu	Phe	Phe
	210		Gln			215					220			-	
Tyr 225	Ala	Asp	Gln	His	His 230	Val	Leu	Leu	Lys	Gly 235	Asp	Leu	Pro	Ile	Leu 240
			Asp	245					250			-		255	
			Ser 260					265			-		270		
		275	His				280					285			
-	290		Trp		-	295	-		-	-	300				-
Ser 305	Val	Tyr	Arg	Leu	Asp 310	His	Ile	Ile	Gly	Phe 315	Phe	Arg	Leu	Trp	Ile 320
Trp			Ser	325		-			330					335	_
Tyr			Gln 340	_				345					350		
		355	Pro				360					365			
_	370		Leu			375				_	380	-			-
385			Asn		390		-			395				-	400
			Leu	405					410					415	
			420					425		Glu		-	430		
-		435	His				440	-				445			
	450		Leu			455					460				
465			Asn		470					475	-				480
			Arg	485	-				490	_				495	-
			Tyr 500					505					510	Ile	His
Lys	Lys	Phe 515	Asn	Gly	Tyr	Ile	Glu 520	Lys	Ile	Leu	Thr	Gly 525	Leu		

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50 60 55 Asn Val Asp Val Leu Ser Leu Phe Ser Asp Val Leu Asp Leu Asp Ala 70 75 Gly Ile Pro Glu Thr Pro Asn Val Leu Leu Ser Asn Glu Met Gln Lys 90 Val Phe Gln Gly Ile Tyr Asn Glu Ile Ser Leu Ile Lys Val Phe Pro 100 105 Asn Gly Asp Lys Ile Val Val Ala Ser Ser Ile Pro Glu His Leu Gly 120 125 Glu Asn Tyr Asn His Lys Ile Asp Ile Pro Lys Asn Thr Pro Phe Leu 135 Ala Ala Leu Lys Gln Ser Pro Lys Asn Gln Glu Val Phe Ser Val Met 150 155 Gln Ala Asn Val Phe Asp Ala Lys Thr Gln Glu Leu Gln Gly Ile Leu 170 165 Tyr Thr Thr Phe Ser Ala Glu Ser Leu Leu Lys Asp Leu Leu Ile Asn 185 Lys Gln Ser Tyr Leu Thr Val Lys Thr Ala Ile Leu Ser Lys Tyr Gly
195 200 205 Val Ile Leu Lys Ala Ser Asp Pro Ala Leu His Leu His Thr Val Tyr

. 215

Pro Asp Met Thr Lys Glu Lys Phe Cys Gln Val Phe Leu Asn Asp Asp 225 230 235 1240

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Pro Cys Pro Ile Asp Ser Glu Leu Gly Pro Leu Thr Leu Ser Pro Leu
245 250 255
Asp Ile Gly Glu Asn Phe Tyr Ser Phe Lys Ile Lys Asp Thr Glu Ile
            260 265
Trp Gly Cys Ile Glu Asn Val Pro Ser Ile Asp Ile Ala Val Leu Ser
275 280 285
Tyr Ala Lys Lys Glu Glu Ser Phe Ala Pro Leu Trp Arg Arg Ala Arg
                       295
                                           300
Met Tyr Thr Ala Tyr Phe Phe Cys Ile Leu Leu Gly Ser Leu Ile Ala
                    310
                                       315
Phe Ile Val Ala Arg Arg Leu Ser Leu Pro Ile Arg Lys Leu Ala Thr
325 330 335
Ala Met Ile Glu Ser Arg Lys Asn Lys Asn Cys Leu Tyr Thr Asp Asp 340 345 350
Ser Leu Gly Phe Glu Ile Asn Arg Leu Gly His Ile Phe Asn Ala Met
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                                                 365
Val Glu Asn Leu His Lys Gln Gln His Leu Ala Lys Thr Asn Phe Glu
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                                            380
Met Lys Glu Asn Ala Gln Asn Ala Leu His Leu Gly Glu Gln Ala Gln 385 $390$
Gln Arg Leu Leu Pro Asn Thr Leu Pro Ser Tyr Pro His Ile Glu Leu 405 410 415
Ala Lys Ala Tyr Ile Pro Ala Ile Thr Val Gly Gly Asp Phe Phe Asp
420 425 430
Val Phe Val Val Gly Glu Gly Ser Lys Ala Arg Leu Phe Leu Ile Val
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Ala Asp Ala Ser Gly Lys Gly Val Asn Ala Cys Gly Tyr Ser Leu Phe
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Leu Lys Asn Met Leu Arg Thr Phe Leu Ser Arg Ser Ser Ser Leu Gln
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Asp Pro Asp Gly Glu Thr Ser Trp Leu Phe His Pro Gly Met Ala Leu 530 540
Gly Phe Leu Pro Glu Val Ala Asn Ile Thr Ser Lys Leu Phe His Pro 545 550 555 560
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Lys Pro Gly Ser Leu Phe Val Leu Tyr Ser Asp Gly Ile Thr Glu Ala
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Ile Gln Gly Leu Thr Gly Lys Ser Ala Ala Asp Ala Val His Arg Leu 595 600 605
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Phe	Gln	Asp	Leu	Phe 165		Asp	Asp	Tyr	Phe 170		Glu	Val	Gln	Leu 175	His
Lys	Met	Ser	Glu 180		Ser	Ile	Ala	Gly 185		Lys	Glu	Glu	Trp	Leu	Lys
Gln	Glu	Tyr 195	Tyr	Ser	Leu	Ile	Glu 200	Lys	Gln	Ile	Lys	Val 205	Asn	Thr	Ala
Val	Leu 210		Ala	Ser	Lys	Arg 215		Gly	Ile	Pro	Thr 220		Ala	Thr	Asn
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Leu	Asn	Val	Gln	Ser 245	Gly	Glu	Thr	Val	Arg 250	Ile	Ala	Lys	Gln	Asn 255	Thr
His	Ile	Pro	Asn 260	Pro	Lys	Arg	Lys	Val 265	Tyr	Arg	Ser	Arg	Glu 270	Tyr	Tyr
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Val	290		Asn			295					300			Thr	
Asp 305	Phe	Ser		_	310	-			-	315				Leu	320
	Leu			Tyr 325					330					Ala 335	
Phe			340				Glu	345			-	-	350	Ser	
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Asp	370					375	Asp			Met	380		Ile		
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Ala				405					410		_			Gly 415	
GTĀ			420				Leu	425					430		Ile
Arg		435					440					445	-	Leu	
	450			-		455		-			460	-		Arg	
465					470					475				Gln	480
Ile		Phe		485			Ala		490					495	Gly
Arg			500					505					510	Lys	
TTE		515					520					525		Asp	
	530					535	Asn				540				Ile
545	Met				550		Gly			Arg 555			Gly		560
ATS.		_		565		-	Gly	-	570					575	
TTE	Cys		580	_			Thr	585		Thr		Gln	590		Met
ьуѕ		595				-	600		-		-	605		Gly	
шуs	Thr 610			Ser		615					620			Lys	-
rnr	стА	GIN	ser	ьeu	Ala	met	ALA	Thr	теп	Pro	ьeu	Asp	Asp	Ala	rnr

625					630					635					640
Thr	Phe	Ser	Leu	Leu 645	His	Gln	Gly	Lys	Thr 650	Met	Gly	Ile	Phe	Gln 655	Met
Glu	Ser	Lys	Gly 660	Met	Gln	Glu	Leu	Ala 665	Lys	Asn	Leu	Arg	Pro 670	Asp	Leu
Phe	Glu	Glu 675	Ile	Ile	Ala	Met	Gly 680	Ala	Leu	Tyr	Arg	Pro 685	Gly	Pro	Met
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Glu 705	Tyr	Asp	His	Pro	Leu 710		Glu	Ser	Ile	Leu 715		Glu	Thr	Tyr	Gly 720
Ile	Met	Val	Tyr	Gln 725	Glu	Gln	Val	Met	Gln 730	Ile	Ala	Gly	Ala	Leu 735	Ala
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		835			Leu		840					845	_		
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865					Phe 870					875					880
_				885	Ile				89Õ	-				895	-
			900		Phe			905					910		
		915			Ser		920					925			
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945					1ys 950	-				955	-				Phe 960
				965	Met				970					975	
			980		Thr			985					990		
		995			Ile		1000)				1005	5		
-	1010)			Arg	1015	5				1020) "			
102	5				Ser 1030)				1035	5			-	1040
	Thr			1045					1050)				1055	5
			1060)	Ile			1065	5				1070)	-
		1075	5		Gln		1080)				1085	5		_
	1090)			Asp	1095	5				1100)			
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Phe Tyr Lys Asp Phe Val Lys Asn Val Asp Ile Glu Leu Leu Asn Gln 305 310 315 320
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Val Thr Tyr Lys Glu Ile Asp Ala Gln Thr Lys Lys Met Lys Thr Asp
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Lys Leu Glu Pro Gly Ser Arg Glu Glu Leu Leu Leu Val Glu Gly Val
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Gln Leu Ser Val Arg Lys Trp Arg His Pro Arg Gly Glu His Tyr 500 505
Gly Asn Val Ile Tyr Ser Asp Thr Glu Leu Asp Arg Ile Gln Met Leu
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Ala Pro Asp Tyr Gly Tyr Gln Gly Ser Trp Thr Leu Val Pro Lys Val
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Gly Ala Gly Gly Lys Val Thr Leu Val Ala Glu Trp Gln Ala Leu Gly
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Tyr Thr Pro Lys Pro Glu Leu Arg Ala Thr Leu Val Pro Asn Ser Leu
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Leu Ile Ser Arg Gly Tyr Ile Val Gly Gly Ser Met Thr Thr Pro Gln
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Asp Tyr Val Val Ser Asp Ile Lys Ser Gln Val Tyr Ala Gly Ser Leu
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Cys Ala Gln Ser Ser Tyr Val Ile Pro Leu His Ser Ser Leu Arg Arg
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His Val Leu Ser Lys Val Leu Pro Glu Leu Pro Gly Glu Thr Pro Leu
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Val Leu His Gly Gln Val Ser Tyr Gly Arg Asn His His Asn Met Thr
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Thr Lys Leu Ala Asn Asn Thr Gln Gly Lys Ser Asp Trp Asp Ser His
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Ser Phe Ala Val Glu Val Gly Gly Ser Leu Pro Val Asp Leu Asn Tyr
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Arg Tyr Leu Thr Ser Tyr Ser Pro Tyr Val Lys Leu Gln Val Val Ser
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Asp Ala Ser His Leu Val Asn Val Ser Ile Pro Met Gly Leu Thr Phe
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Lys His Glu Ser Ala Lys Pro Pro Ser Ala Leu Leu Leu Thr Leu Gly
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Tyr Ala Val Asp Ala Tyr Arg Asp His Pro His Cys Leu Thr Ser Leu
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Asp Cys Phe Ala Ser Gly Ser Cys Glu Leu Arg Ser Ser Ser Arg Ser
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atgati gati tcto gaaa	tteed ceeti acto acco	ata ccg ag	aaaga aatta aacto tagga aaaao	agaco ctaco attat	et a	aaaat aaaaq ccaga	tacga gtgtd actct	a gta c tca c cc	aaaca actaa tgtt	aagg attc cttg	tagg aaaa	agta gacto atta	aac gct aga	tttci tcaa ggata	ttaaa caaat aacag	it :a ;t	60 120 180 240 300 306
<213 <213	0> 5! 1> 72 2> D1 3> C	29 NA	achor	natis	5 D :	sero	7ar										
gtgg gaca gggac gggac ggaca cgtg tctt taca ctta aagga <210 <211	ccaa atagt gacet gacet ataat gaatt tttt gaaaa aaaga ctaac acaac 1> 28	gca att at aa gta aa ct aa gca aa ag ag aa gca ag ag ag ag ag ag ag ag ag ag ag ag ag	acttotactactactactactactactactactactactactact	ttete ateaa eetaa eeceae etgtt eegta eegta aata	gg at a control at	ttggggatti ttttt ttatt agcet actec ggaaa tegac acaaa aatgt	ggget caaaa catee cagga cagaa cagaa cataac cataac cataac catat	agi a to gaa gaa gaa ga co gaa co caa co caa co caa co caa co caa co caa co caa co caa co caa co caa co caa caa	tgto aato acagi tgagi gtta gtta gttti agaa aatgi aatgi	agaa attt ttta tact acga ccta gaac tata gata	ataa gega gaga gega aaga ttet acat taga ttto	accas aacca actect accet agge teet acate eccec	aaa gga ttt ttt gtt gtt cag	agger aaaaq tatte ttate tgtte acaaa aatae cgagr ccgtt agata	tgca gatag gacat gacat gcagg aagat gcttt cctat	ic it ia ia ia ia ia is ia is is is is is is is is is is is is is	60 120 180 240 300 360 420 480 540 600 720 729
	3> C. D> 56		acnon	atis	3 D 3	serov	ar										
			Gln	His 5	Lys	Lys	Ile	Ser	Glu 10	Glu	Thr	Ile	Ala	Cys 15	Asp		
			Arg 20					25					30				
		35	Thr				40	_		-		45					
-	50		Pro			55					60						
Pro	Gln	Ile	His	Thr	Ser	Ile	Ile	Asp	Phe	Lys	Leu	Gly	Ser	Pro	Gly		

Ala Ala Leu Thr Val Asp Leu Cys Ser Phe Leu Pro Asn Ala Thr Ala 90 Ala Ile Met Leu Gly Met Cys Gly Gly Leu Arg Ser His Tyr Gln Ile 100 105 Gly Asp Tyr Phe Val Pro Val Ala Ser Ile Arg Lys Asp Gly Thr Ser 115 120 125 Asp Ala Tyr Phe Pro Pro Glu Val Pro Ala Leu Ala Asn Phe Val Val 130 135 140 Gln Lys Met Ile Thr Asn Ile Leu Glu Ala Lys Asn Leu Pro Tyr His 145 150 155 160 Ile Gly Ile Thr His Thr Thr Asn Ile Arg Phe Trp Glu Phe Asn Lys $165 \hspace{1.5cm} 170 \hspace{1.5cm} 175$ Glu Phe Arg Arg Lys Leu Tyr Glu Asn Lys Ala Gln Thr Val Glu Met 180 185 190 Glu Cys Ala Thr Leu Phe Ala Ala Gly Tyr Arg Arg Asn Leu Pro Leu 195 200 205 Gly Ala Leu Leu Leu Ile Ser Asp Leu Pro Leu Arg Lys Asp Gly Ile 210 215 220 Lys Thr Lys Glu Ser Ser Ser Ala Val Leu Asn Ser His Thr Lys Glu 230 235 His Ile Leu Thr Gly Val Glu Val Phe Ala Ser Leu Gln Glu Lys Ser 245 250 255 Gly Pro Gly Ile Lys Lys Thr Lys Gly Leu Pro His Met Glu Phe Gly 260 265 270 Gln Ala Asp Asp Ser Leu Ser Glu Gln Thr Glu Val Ser Gly Gly Asp 280 Phe

<210> 561 <211> 394 <212> PRT <213> C. Trachomatis D serovar

<400> 561 Met Ser Lys Glu Thr Phe Gln Arg Asn Lys Pro His Ile Asn Ile Gly
1 5 10 15 Thr Ile Gly His Val Asp His Gly Lys Thr Thr Leu Thr Ala Ala Ile 20 25 30 Thr Arg Ala Leu Ser Gly Asp Gly Leu Ala Asp Phe Arg Asp Tyr Ser 40 Ser Ile Asp Asn Thr Pro Glu Glu Lys Ala Arg Gly Ile Thr Ile Asn 55 Ala Ser His Val Glu Tyr Glu Thr Ala Asn Arg His Tyr Ala His Val 75 70 Asp Cys Pro Gly His Ala Asp Tyr Val Lys Asn Met Ile Thr Gly Ala 85 90 95 Ala Gln Met Asp Gly Ala Ile Leu Val Val Ser Ala Thr Asp Gly Ala Met Pro Gln Thr Lys Glu His Ile Leu Leu Ala Arg Gln Val Gly Val Pro Tyr Ile Val Val Phe Leu Asn Lys Ile Asp Met Ile Ser Glu Glu 130 135 140 Asp Ala Glu Leu Val Asp Leu Val Glu Met Glu Leu Val Glu Leu Leu 145 150 150 160 Glu Glu Lys Gly Tyr Lys Gly Cys Pro Ile Ile Arg Gly Ser Ala Leu 165 170 175 Lys Ala Leu Glu Gly Asp Ala Ala Tyr Ile Glu Lys Val Arg Glu Leu 180 185 190 Met Gln Ala Val Asp Asp Asn Ile Pro Thr Pro Glu Arg Glu Ile Asp 195 200 205

Lys Pro Phe Leu Met Pro Ile Glu Asp Val Phe Ser Ile Ser Gly Arg 210 215 220 Gly Thr Val Val Thr Gly Arg Ile Glu Arg Gly Ile Val Lys Val Ser 230 235 Asp Lys Val Gln Leu Val Gly Leu Arg Asp Thr Lys Glu Thr Ile Val 245 250 Thr Gly Val Glu Met Phe Arg Lys Glu Leu Pro Glu Gly Arg Ala Gly 265 270 260 Glu Asn Val Gly Leu Leu Leu Arg Gly Ile Gly Lys Asn Asp Val Glu 280 Arg Gly Met Val Val Cys Leu Pro Asn Ser Val Lys Pro His Thr Gln 290 295 300 Phe Lys Cys Ala Val Tyr Val Leu Gln Lys Glu Glu Gly Gly Arg His 310 315 Lys Pro Phe Phe Thr Gly Tyr Arg Pro Gln Phe Phe Phe Arg Thr Thr 325 330 335 Asp Val Thr Gly Val Val Thr Leu Pro Glu Gly Ile Glu Met Val Met 340 345 Pro Gly Asp Asn Val Glu Phe Glu Val Gln Leu Ile Ser Pro Val Ala 355 360 365 Leu Glu Glu Gly Met Arg Phe Ala Ile Arg Glu Gly Gly Arg Thr Ile 375 380 Gly Ala Gly Thr Ile Ser Lys Ile Ile Ala 390 <210> 562 <211> 550 <212> PRT <213> C. Trachomatis D serovar <400> 562 Met Glu Ser Ser Arg Ile Leu Ile Thr Ser Ala Leu Pro Tyr Ala Asn 10 Gly Pro Leu His Phe Gly His Ile Thr Gly Ala Tyr Leu Pro Ala Asp 20 25 Val Tyr Ala Arg Phe Gln Arg Leu Gln Gly Lys Glu Val Leu Tyr Ile 40 4.5 Cys Gly Ser Asp Glu Tyr Gly Ile Ala Ile Thr Leu Asn Ala Glu Leu 60 55 Ala Gly Met Gly Tyr Gln Glu Tyr Val Asp Met Tyr His Lys Leu His 75 Lys Asp Thr Phe Lys Lys Leu Gly Ile Ser Val Asp Phe Phe Ser Arg 85 90 Thr Thr Asn Thr Tyr His Pro Ala Ile Val Gln Asp Phe Tyr Arg Asn 105 Leu Gln Glu Arg Gly Leu Val Glu Asn Gln Val Thr Glu Gln Leu Tyr 120 125 Ser Glu Glu Glu Gly Lys Phe Leu Ala Asp Arg Tyr Val Val Gly Thr 130 135 140 Cys Pro Lys Cys Gly Phe Asp Arg Ala Arg Gly Asp Glu Cys Gln Gln 145 150 155 Cys Gly Ala Asp Tyr Glu Ala Arg Asp Leu Lys Glu Pro Arg Ser Lys 165 170 175 Leu Thr Gly Ala Ala Leu Ser Leu Arg Asp Thr Glu His Ala Tyr Leu 180 185 190 His Leu Glu Arg Met Lys Glu Asp Leu Leu Ala Phe Val Gln Gly Ile 200 Tyr Leu Arg Pro His Met Arg Asn Phe Val Thr Asp Tyr Ile Glu His 210 . 215 220 Leu Arg Fro Arg Ala Val Thr Arg Asp Leu Ser Trp Gly Ile Fro Val 225 230 235 240

Pro Asp Leu Glu Asn Lys Val Phe Tyr Val Trp Phe Asp Ala Pro Ile 245 250 255 Gly Tyr Ile Ser Gly Thr Met Asp Trp Ala Ala Ser Ile Gly Asp Pro 260 265 270 Glu Ala Trp Lys Lys Phe Trp Leu Asp Asp Thr Val Thr Tyr Ala Gln 275 280 285 Phe Ile Gly Lys Asp Asn Thr Ser Phe His Ala Ala Ile Phe Pro Ala 290 295 300 Met Glu Ile Gly Gln Ser Leu Pro Tyr Lys Lys Val Asp Ala Leu Val 310 315 Thr Ser Glu Phe Leu Leu Glu Gly Phe Gln Phe Ser Lys Ser Asp 330 325 Gly Asn Phe Ile Asp Met Asp Ala Phe Leu Glu Thr Tyr Ser Leu Asp 340 345 350Lys Leu Arg Tyr Val Leu Ala Ala Ile Ala Pro Glu Thr Ser Asp Ser 355 360 365 Glu Phe Ser Phe Gln Glu Phe Lys Thr Arg Cys Asn Ser Glu Leu Val Gly Lys Tyr Gly Asn Phe Val Asn Arg Val Leu Ala Phe Ala Val Lys 385 390 395 400 Asn Gly Cys Thr Glu Leu Ser Ser Pro Gln Leu Glu Gln Lys Asp Leu
405 410 415 Asp Phe Ile Ser Lys Ser Gln Lys Leu Ala Lys Asp Ala Ala Glu His 420 425 430Tyr Ala Gln Tyr Ser Leu Arg Lys Ala Cys Ser Thr Ile Met Glu Leu 435 440 445 Ala Ala Leu Gly Asn Gly Tyr Phe Asn Asp Glu Ala Pro Trp Lys Leu 450 455 Ala Lys Glu Gly Asn Trp Asn Arg Val Arg Ala Ile Leu Phe Cys Ala 470 475 Cys Tyr Cys Gln Lys Leu Leu Ala Leu Ile Ser Tyr Pro Ile Met Pro 485 490 Glu Thr Ala Leu Lys Ile Leu Glu Met Ile Ala Pro His Ser Leu Asp $500 \hspace{1cm} 505 \hspace{1cm} 510$ Leu Gly Ser Gln Asp Pro Asp Arg Leu Gln Ser Leu Trp Thr Asp Ser 515 520 525 Phe Phe Asp Tyr Ser Glu Glu Lys Phe Ser Leu Lys Glu Pro Glu Leu 530 535 540 Leu Phe Thr Met Val Glu <210> 563 <211> 100 <212> PRT <213> C. Trachomatis D serovar <400> 563

<210> 564

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<213> C. Trachomatis D serovar
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Phe Pro Lys Leu Phe Ile Ile Ser Ala Pro Ala Gly Ala Gly Lys Thr
                                25
Thr Leu Thr His Met Leu Gln Arg Glu Phe Pro Asp Ala Phe Glu Lys
                            40
Thr Val Ser Ser Thr Thr Arg Ser Ala Arg Pro Gly Glu Val His Gly
                        55
Val Asp Tyr Leu Phe Val Ser Glu Asp Asp Phe Lys Gln Ser Leu Asp
65 70 75 80
Arg Glu Asp Fhe Leu Glu Trp Val Phe Leu Phe Gly Thr Tyr Tyr Gly
                85
Thr Ser Lys Ala Glu Ile Ser Arg Val Leu Gln Lys Gly Lys His Cys
           100
                                 105
Ile Ala Val Ile Asp Val Gln Gly Ala Leu Ala Leu Lys Lys Gln Met
                            120
                                                 125
Pro Ala Val Thr Ile Phe Ile Gln Ala Pro Ser Gln Glu Glu Leu Glu
                       135
Arg Arg Leu Asn Ala Arg Asp Ser Glu Lys Asp Phe Gln Lys Lys Glu
145 150 155 160
Arg Leu Glu His Ser Ala Val Glu Ile Ala Ala Ala Ser Glu Phe Asp
165 170
Tyr Val Val Val Asn Asp Asp Leu Ile Thr Ala Tyr Gln Val Leu Arg 180 185 190
Ser Ile Phe Ile Ala Glu Glu His Arg Met Ser His Gly
<210> 565
<211> 602
<212> PRT
<213> C. Trachomatis D serovar
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Met Lys Pro Tyr Lys Ile Glu Asn Ile Arg Asn Phe Ser Ile Ile Ala
                                    10
His Ile Asp His Gly Lys Ser Thr Ile Ala Asp Arg Leu Leu Glu Ser
                                 25
Thr Ser Thr Ile Glu Gln Arg Glu Met Arg Glu Gln Leu Leu Asp Ser
                             40
Met Asp Leu Glu Arg Glu Arg Gly Ile Thr Ile Lys Ala His Pro Val
                        55
Thr Met Thr Tyr Glu Tyr Glu Gly Glu Thr Tyr Glu Leu Asn Leu 11e
65 70 75 80
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Asp Thr Pro Gly His Val Asp Phe Ser Tyr Glu Val Ser Arg Ser Leu

Glu Ile Ile Pro Val Leu Asn Lys Ile Asp Leu Pro Ala Ala Gln Pro

Glu Ala Ile Lys Lys Gln Ile Glu Glu Phe Ile Gly Leu Asp Thr Ser

Asn Thr Ile Ala Cys Ser Ala Lys Thr Gly Gln Gly Ile Pro Glu Ile

135

150

90 Ala Ala Cys Glu Gly Ala Leu Leu Ile Val Asp Ala Ala Gln Gly Val

155

125

140

105 Gln Ala Gln Ser Leu Ala Asn Val Tyr Leu Ala Leu Glu Arg Asp Leu 120

85

				165					170					175	
Leu	Glu	Ser	Ile 180	Ile	Arg	Leu	Val	Pro 185	Pro	Pro	Lys	Pro	Pro 190	Gln	Glu
	Glu	195					200					205			
	11e 210					215				-	220		-	-	_
225	Arg				230					235					240
_	Ile	-		245					250				_	255	
	Ala		260			-		265				_	270		-
_	Val	275		-			280				-	285			-
	Pro 290					295			_		300				_
305	Tyr				310					315					320
_	Arg			325		-			330					335	
	His		340					345					350		
	Leu	355					360			-		365	-		-
	370					375					380			_	
385	Lys				390	-			'	395			-		400
	Ile			405				-	410					415	
	Gln		420					425					430		
	Ile	435					440					445			
	Tyr 450					455					460			-	-
465	Lys				470	_	_	_		475	-	-	-		480
_	Tyr	-	-	485				-	490					495	-
	Ala		500				-	505			-	-	510		
	Lys	515					520					525			
	Leu 530					535					540	_	-		
545	Arg				550					555				-	560
	Gly			565					570					575	
	Gly		580			-		585		Lys	Val	Ser	Ile 590	Pro	Asn
Thr	Ala	Phe 595	Val	Glu	Val	Leu	Lys 600	Met	Glu						

<210> 566 <211> 324 <212> PRT <213> C. Trachomatis D serovar

<400> 566 Met Glu Leu Leu Pro His Glu Lys Gln Val Val Glu Tyr Glu Lys Thr 5 10 Ile Ala Glu Phe Lys Glu Lys Asn Lys Glu Asn Ser Leu Leu Ser Ser 20 25 Ser Glu Ile Gln Lys Leu Asp Lys Arg Leu Asp Arg Leu Lys Glu Lys 40 Ile Tyr Ser Asp Leu Thr Pro Trp Glu Arg Val Gln Ile Cys Arg His 50 Pro Ser Arg Pro Arg Thr Val Asn Tyr Ile Glu Gly Met Cys Glu Glu 65 70 75 80 Phe Val Glu Leu Cys Gly Asp Arg Thr Phe Arg Asp Asp Pro Ala Val 85 90 95 Val Gly Gly Phe Ala Lys Ile Gln Gly Gln Arg Phe Met Leu Ile Gly 105 100 Gln Glu Lys Gly Cys Asp Thr Lys Ser Arg Met His Arg Asn Phe Gly 115 120 125 Met Leu Cys Pro Glu Gly Phe Arg Lys Ala Leu Arg Leu Ala Lys Met 135 140 Ala Glu Lys Phe Gly Leu Pro Ile Ile Phe Leu Val Asp Thr Pro Gly 150 155 Ala Phe Pro Gly Leu Thr Ala Glu Glu Arg Gly Gln Gly Trp Ala Ile 165 170 Ala Thr Asn Leu Phe Glu Leu Ala Arg Leu Ala Thr Pro Ile Ile Val 180 185 Ile Val Ile Gly Glu Gly Cys Ser Gly Gly Ala Leu Gly Met Ala Ile 195 200 205 Gly Asp Val Val Ala Met Leu Glu His Ser Tyr Tyr Ser Val Ile Ser 210 $\,$ 220 $\,$ Pro Glu Gly Cys Ala Ser Ile Leu Trp Lys Asp Pro Lys Lys Asn Ser 230 235 Asp Ala Ala Met Leu Lys Met His Gly Glu Asp Leu Lys Gly Phe 245 250 255 Ala Ile Val Asp Ala Val Ile Lys Glu Pro Ile Gly Gly Ala His His 260 265 270 Asn Pro Ala Ala Thr Tyr Arg Ser Val Gln Glu Tyr Val Leu Gln Glu 275 280 285 Trp Leu Lys Leu Lys Asp Leu Pro Val Glu Glu Leu Leu Glu Lys Arg 290 295 300 Tyr Gln Lys Phe Arg Thr Ile Gly Leu Tyr Glu Thr Ser Ser Glu Ser 305 310 315 Asp Ser Glu Ala

<211> 646 <212> PRT

<213> C. Trachomatis D serovar

<400> 567

<210> 567

Met Lys Leu Leu Lys Ala Ile Leu Arg His Lys Lys His Leu Val 10 Leu Phe Gly Phe Ser Leu Leu Ser Ile Leu Gly Leu Thr Ile Thr Ser Gln Ala Glu Ile Phe Ser Leu Gly Leu Ile Ala Lys Thr Gly Pro Asp 40 Thr Phe Leu Leu Phe Gly Lys Gln Glu Gly Ala Ser Leu Val Lys Arg 60 55 Lys Glu Leu Ser Lys Asp Gln Leu Leu Glu Gln Trp Asp Asn Ile Val 65 70 75 80 Gly Glu Gly Asp Thr Leu Ser Leu Pro Gln Ala Asn Ala Tyr Ile Ala

Lys	His	Ser	Gly 100	85 Gly	Ser	Gln	Ser	Ile 105	90 Thr	Lys	Arg	Leu	Ser 110	95 Ala	Tyr
Leu	Ser	Gly 115		Phe	Asp	Phe	Ser 120		Leu	Gln	Cys	Leu 125		Leu	Phe
Leu	Val 130		Val	Ala	Ile	Leu 135		Ser	Thr	Thr	Leu 140		Phe	Gl n	Arg
Phe 145	Leu	Ala	Gln	Leu	11e 150	Ala	Ile	Arg	Val	Ser 155	Cys	Ser	Leu	Arg	Lys 160
	Tyr			165					170					175	
	Asp		180					185				-	190		
	Ala	195					200					205			
	Thr 210					215			-		220			•	-
225	Cys		-		230		Ala			235					240
vai	Ile			245			Ala		250		_			255	-
Len	Thr		260					265		-			270	_	
	Gln	275					280					285		_	-
-	290 Ser	-		-		295		-			300				
305	Ala				310		Gly			315					320
Glu				325			Gly		330					335	
Ile	Lys	Lys	340				Asn	345				Trp	350		
Ala	Ala	355 Glu	Arg	Phe	Tyr	Glu	360 Val	Leu	Asp	Leu	Ala	365 Lys	Gln	Gln	Ser
	370 Val	Ser	Glu	Lys		375 Asn	Glu	Phe	Gln		380 Leu	Gln	His	Ser	
385 Gln	Phe	Cys	Asn		390 Ser	Phe	Gly	Tyr	Val 410	395 Glu	Asp	Ser	Pro	Val 415	400 Leu
Ser	Asp	Phe	Asn 420	405 Leu	Val	Leu	Lys	Lys 425		Glu	Ala	Ile	Gly 430		Val
Gly	Pro	Thr 435		Ser	Gly	Lys	Ser 440		Ile	Ala	Lys	Leu 445		Pro	Arg
Leu	Tyr 450		Val	Ser	His	Gly 455	Glu	Leu	Leu	Ile	Asp 460		Leu	Pro	Ile
Arg 465	Ser	Tyr	Суз	Ьуз	Asn 470		Leu	Arg	Ьуз	His 475		Gly	Ċйз	Val	Leu 480
Gln	His	Pro	Phe	Leu 485	Phe	Tyr	Asp	Thr	Val 490	Trp	Asn	Asn	Leu	Thr 495	Суз
Gly	Arg	Thr	Phe 500	Ser	Glu	Glu	Glu	Val 505	Phe	His	Ala	Leu	Lys 510	Gln	Ala
	Ala	515					520				_	525			
	Glu 530					535					540			_	
545	Ile				550					555					560
GLu	Ala	Thr	Ser	Ala 565	Leu	Asp	Ala	Ile	Ser 570	Glu	Asn	Tyr	Val	Lys 575	G1u

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Ile Val Gly Gln Leu Lys Gly Arg Cys Thr Gln Ile Ile Ile Ala His 580 \hspace{1cm} 585 \hspace{1cm} 585 \hspace{1cm} 590
Lys Leu Ser Thr Leu Glu Tyr Val Asp Arg Ile Val Tyr Leu Glu Gln
595 600 605
Gly Lys Lys Ile Ala Glu Gly Thr Lys Glu Glu Leu Leu Asp Ser Cys
                        615
Pro Ala Phe Gln Arg Met Trp Val Leu Ser Gly Ala Lys Asp Trp Glu
                                          635
                     630
Leu Asn Ala Val Val Lys
<210> 568
<211> 414
<212> PRT
<213> C. Trachomatis D serovar
<400> 568
Met Phe Ser Ser Ala Ile Val Ile Leu Thr Ala Ile Phe Val Leu Cys 1 \hspace{1cm} 5 \hspace{1cm} 10 \hspace{1cm} 15
Ser Gly Phe Val Ser Leu Ser His Ile Ala Leu Phe Ser Leu Pro Ser
   20
                                  25
Ser Leu Ile Ala His Tyr Ser His Ser Lys Asn Arg Gln Leu Arg Gln
 35
                           40
Ile Ala Asn Leu Met Ala Tyr Pro Asn His Leu Leu Met Thr Leu Val
Phe Phe Asp Ile Gly Ile Asn Ile Gly Val Gln Asn Cys Ile Ala Thr
Leu Val Gly Asp Ser Ala Ser Leu Leu Leu Thr Val Gly Val Pro Leu
                 85
                                       90
Pro Tyr Asn Ala Arg Ile Ala Lys Ile Val Thr Pro Ile Ile Phe Ala
Ser Thr Lys Ser Phe Arg Pro Ile Phe Asp Trp Ala Ile Ser Gly Ile 130 $135\ 
Asn Phe Ile Val Gln Lys Met Leu Ala Arg Gln Glu Ser Asp Phe Ile
145 150 155 160
Gln Pro Gln Glu Leu Lys Glu Val Leu Arg Ser Cys Lys Asp Phe Gly 165 170 175
Val Val Asn His Glu Glu Ser Arg Leu Leu Phe Gly Tyr Leu Ser Met
180 185 190
Glu Glu Gly Ser Ile Lys Glu Arg Met Thr Pro Lys Gln Glu Ile Ile
195 200 205
Phe Tyr Asp Val Leu Thr Pro Ile Glu Asn Leu Tyr Lys Leu Phe Ser 210 225
Gly Pro Lys Gln Ser Tyr Ser Lys Val Leu Val Cys Lys Gly Gly Leu 225 230 235 240
Gln Asn Leu Leu Gly Val Cys Ser Ala Lys Leu Leu Leu Leu Tyr Lys
245 250 255
Glu Lys Leu Gln Ser Ala Glu Glu Leu Leu Pro Leu Leu Arg Lys Pro 260 \hspace{1.5cm} 265 \hspace{1.5cm} 270 \hspace{1.5cm}
His Tyr Ile Pro Glu Thr Val Ser Ala Lys Thr Ala Leu Tyr His Leu 275 280 285
Ala Gly Glu Asp Cys Gly Leu Gly Ile Ile Ile Asp Glu Tyr Gly Ser 290 295 300
Ile Glu Gly Leu Ile Thr Gln Asn Asp Leu Phe Lys Ile Val Ser Asp
Gly Val Ala His Asn Arg Pro Ser Phe Lys Gln Phe Ala His Ser Asp
                325 330 335
Lys Asn Val Val Ile Ala Ala Gly Thr Tyr Glu Leu Ser Asp Phe Tyr 340 345 350
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Asp Leu Phe Gly Val Asp Leu Pro Thr Thr Ala Asn Cys Val Thr Ile 355 360 365
Gly Gly Trp Leu Thr Glu Gln Leu Gly Glu Ile Pro Glu Thr Gly Thr 370 375 380
Lys Phe Ala Trp Gly Gln Phe Val Phe Gln Ile Leu Asp Ala Ala Pro
                    390
                              395
Asn Cys Val Lys Arg Val Tyr Ile Arg Lys Thr His Gly Asn
                405
                             410
<210> 569
<211> 404
<212> PRT
<213> C. Trachomatis D serovar
<400> 569
Met Glu Thr Asn Ser Pro Phe Phe Trp Leu Gly Val Asn Leu Leu Cys
Ile Phe Val Gln Gly Phe Phe Ser Met Met Glu Met Ala Cys Ile Ser 25 30
Phe Asn Arg Val Arg Leu Gln Tyr Tyr Leu Thr Lys Ser Asn Lys Lys
35 40 45
Ala Ser Tyr Ile Asn Phe Leu Val Arg Arg Pro Tyr Arg Leu Phe Gly
                     55
Thr Val Met Leu Gly Val Asn Ile Ala Leu Gln Ile Gly Ser Glu Ser
Ser Arg Thr Cys Tyr Lys Leu Leu Gly Ile Ser Pro Glu Tyr Ala Pro
85 90 95
Ala Thr Gln Ile Ile Leu Val Val Ile Phe Ala Glu Leu Ile Pro Leu
100 105 110
Ala Ile Ser Arg Lys Ile Pro Glu Lys Ile Ala Leu Lys Gly Ala Pro
115 120 125
Ile Leu Tyr Phe Ala His Tyr Leu Phe Tyr Pro Leu Ile Gln Cys Val
130 135 140
Gly Gly Ile Thr Asn Met Ile Tyr Phe Ile Leu Asn Ile Lys Glu Glu
145 150 155 160
Thr Leu His Ser Thr Leu Ser Arg Asp Glu Leu Gln Lys Thr Leu Glu
165 170 175
Thr His His Glu Glu His Asp Phe Asn Val Ile Ala Thr Asn Ile Phe
         180
                           185 190
Ser Leu Ser Ala Thr Ser Val Glu Gln Val Cys Gln Tyr Leu Asp Gln
195 200 205
Ile Pro Ile Leu Ser Ala Thr Ala Ser Val Arg Asp Val Cys Gln Leu
210 215 220
Val Arg Arg His Arg Leu Asp Phe Val Pro Val Tyr His Lys Val Lys
                     230
                                         235
Lys Asn Val Val Gly Ile Ala Phe Pro Lys Asn Leu Ile Asn Arg Asn 245 250 255
Pro Ser Asp Pro Val Val Pro Tyr Leu Ser Ser Pro Trp Phe Ile Thr 260 265
Ala Lys Ser Lys Leu Ile His Ala Ile Gln Glu Phe Arg Lys Asn Ser
275 280 285
Ser Asn Val Ala Ile Val Leu Asn Asn Gly Glu Pro Met Gly Val
290 295 300
Leu Gly Leu His Thr Val Phe Lys Thr Leu Phe Asn Thr Arg Asn Ile
305 310 315
Ala Gln Leu Lys Pro Lys Pro Thr Ser Leu Ile Glu Arg Thr Phe Ser
                325 330 335
Gly Asn Thr Pro Leu Ser Glu Ile Glu Asn Glu Leu Asp Ile Ile Phe
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340 345 350 Met Asp Asp Cys Thr Thr Ile Glu Gln Leu Met Leu Lys Leu Leu 355 360 365

Asp Thr Pro Pro Glu Val Gly Ala Ser Ile Ile Ile Asn Asp Leu Leu 375 Leu Glu Val Lys Glu Ile Ser Leu Tyr Gly Ile Lys Thr Val Ala Ile 390 Lvs Asp Thr Leu <210> 570 <211> 539 <212> PRT <213> C. Trachomatis D serovar <400> 570 Met Cys Cys Val Asp Gly Ser Asn Ser Ile Gln Gln Arg Met Arg Phe 10 Cys Glu Tyr Arg Thr Ala Ala Gln Glu Ala Lys Thr Ser Leu Ser Ser Asp Cys Ser Leu Leu Glu Ala Arg Leu Ala Leu Arg Ala Leu Ala Lys 35 40 45 40 His His Glu Tyr Ser Ala Trp Arg Glu Ala Phe Leu Arg Ser Gln Glu
50 60 Arg Phe Pro Ser Leu Glu Ala Asp Arg Asp Ile His Glu Asp Leu Ala 65 70 75 80 Ala Ser Leu Leu Gln Lys Asn Ile Arg His Ser Ser Leu Thr Val Arg 90 Pro Ile Val Leu Gln Ala Leu Ser Asp Asp Ser Asp Thr Val Arg Glu 120 115 125 Ile Ala Val Gln Val Ala Val Met Tyr Gly Ser Ser Cys Leu Leu Arg 135 140 Ala Val Gly Asp Leu Ala Lys Asn Asp Ser Ser Ile Gln Val Arg Ile 145 150 155 160 Thr Ala Tyr Arg Ala Ala Ala Val Leu Glu Ile Gln Asp Leu Val Pro 165 170 175 His Leu Arg Val Val Val Gln Asn Thr Gln Leu Asp Gly Thr Glu Arg Arg Glu Ala Trp Arg Ser Leu Cys Val Leu Thr Arg Pro His Ser Gly 195 200 205 Val Leu Thr Gly Ile Asp Gln Ala Leu Met Thr Cys Glu Met Leu Lys 215 220 Glu Tyr Pro Glu Lys Cys Thr Glu Glu Gln Ile Arg Thr Leu Leu Ala 225 230 235 240 Ala Asp His Pro Glu Val Gln Val Ala Thr Leu Gln Ile Ile Leu Arg 245 250 255 Gly Gly Arg Val Phe Arg Ser Ser Ser Ile Met Glu Ser Val Gln Lys 260 265 270 Leu Ala Cys Asn Ser Leu Ser Ala Arg Val Gln Met Gln Ala Ala Ala 275 280 285 Ile Leu Tyr Leu Glu Gly Asp Pro Phe Gly Glu Asp Lys Leu Thr Glu 290 295 300 Gly Leu Ser Ala Thr Ser Ser Ile Leu Cys Glu Ala Ala Ser Glu Ala 305 310 315 320

Val Cys Ser Leu Gly Ile His Gly Val His Leu Ala Gly Arg Phe Leu 325 330 335 Ser Lys Val Gln Gly Met Arg Ser Arg Val Asn Leu Ala Phe Ala Leu 340 350 Leu Val Ser Arg Glu Lys Val Glu Glu Ala Gly Asp Val Val Ala Ser 360

Phe Ile His Arg Ile Glu Pro Cys Arg Ala Ile Glu Gln Phe Leu Cys $370 \hspace{1cm} 375 \hspace{1cm} 380$

365

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Glu Asp Gln Lys Ile Phe Val Ala Ser Ser Pro Leu Gln Val Glu Ile
                           395
                  390
Met Lys Arg Asp Leu Ala Lys Lys Ile Ile Arg Leu Leu Val Ala Ala
              405
                                 410
Gln Tyr Ser Lys Ala Lys Met Val Val Ala Gln Tyr Leu Ala Gly Gln
           420
                          425
Gln Val Gly Trp Ser Phe Cys Ser Glu Val Phe Trp Glu Glu Gly Asp
       435
                         440
Ser Glu Asp Phe Val Glu Pro Leu Gln Glu Glu Ser Phe Ala Phe Ala
                    455
                                        460
Leu Glu Lys Ala Leu Ser Phe Leu Gln Arg Glu Gly Gly Glu Ala Gly
                 470
                                     475
Leu His Ala Val Ile Ser Leu Tyr Pro His Ser Arg Trp Gln Asp Lys
              485
                                490
Leu Thr Ile Leu Glu Ala Ile Ala Tyr Ser Glu Asn Arg Ile Ala Thr
           500
                             505
                                                510
Cys Phe Leu Arg Glu Arg Cys Leu Gln Glu Ala Ala Ser Leu Gln Ser
                        520
Ala Ala Ala Gly Ala Val Phe Ala Leu Phe Lys
                     535
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<210> 571 <211> 104 <212> PRT

<213> C. Trachomatis D serovar

<400> 571

Met Gln Thr Ser Arg Ile Ser Ser Phe Phe Arg Gly Leu Val His Leu I 1 5 10 15 Tyr Arg Trp Ala Ile Ser Pro Phe Leu Gly Ala Pro Cys Arg Phe Phe 20 25 30 Pro Thr Cys Ser Glu Tyr Ala Leu Val Ala Leu Lys Lys His Pro Leu 40 Arg Lys Ser Leu Phe Leu Ile Ala Lys Arg Leu Leu Lys Cys Gly Pro 50 60 Fro Cys Ile Gly Gly Ile Asp Leu Val Pro Arg Thr Ser Val Glu Glu Glu 65 70 75 86 Pro Thr Pro Leu Ala Glu Ser Pro Asp Asp Arg Thr Ser Val Pro His Thr Gln Glu Thr Ser Val Pro His Thr Gln Glu Thr Ser

<210> 572

<211> 336 <212> PRT

<213> C. Trachomatis D serovar

<400> 572

Met Gln Leu Phe Phe Gly Arg Phe Tyr Glu Val Ala Cys Ile Val Ala 10 Ser Ile Leu Arg Glu Arg Asp Val Gly Val Phe Met Gly Ile Glu Gly 25 30 Arg Gly Ser Gly Ala Met Gln Ser Lys Lys Thr Ile Lys Trp Leu Lys 35 40 45 Gln Ala Leu Val Leu Ser Ser Ile Val Asn Ile Leu Leu Leu Leu Leu 5.5 Ile Tyr Ser Thr Val Phe Arg Lys Asp Ile Tyr Lys Leu Arg Val Phe 75 70 Pro Gly Asn Leu Ile Ala Lys Ser Ser Arg Ile Gly Lys Ile Pro Glu 90

Asp Ile Leu Glu Arg Leu Glu Asn Ala Ser Phe Ala Asp Leu Leu Ala

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105
Leu Leu Gln Glu Glu Arg Met Val Phe Gly His Pro Leu Lys Ser Trp
115 120 125
Ala Leu Gly Val Ser Ile Gln Lys Tyr Phe Val Asp Ile Ala Pro Met
                   135
                               140
Leu Thr His Pro Leu Thr Phe Ile Arg Leu Lys Ser Pro Glu Arg Thr
       150 155
Trp Leu Leu Pro Asp Ile Asn Asp Gln Glu Phe Thr Arg Ile Cys Gln
165 170 175
Tyr Leu Leu Thr Glu Arg Phe Pro Phe Ser Ser Arg Gly Phe Phe Arg
          180
                           185
Ile Met Val Arg Asp Cys Glu Ala Gly Met Val Asp Glu Asp Val Leu
195 200 205
Tyr Arg Phe Cys His Leu Fro Glu Phe Leu Tyr Val Arg Ser Leu Leu
210 215 220
Phe Gly Ala Glu Ile Glu Ala Ala Ser Val Ala Ser Leu Ala Arg Met
               230
                                 235 240
Ile Ile Gln Gly Gly Glu Asp Leu Phe Phe Ser Leu Cys Cys Leu Glu
245 250 255
Asn Arg Gln Thr Ala Ile Ser Asp His Gln Arg Arg Cys Phe Leu Lys 260 265 270
Ala Tyr Val Asp Arg Gln Glu Pro Leu Ala Ala Leu Leu Leu Val
    275 280 285
His Asp Ala Asp Trp Val Leu His Glu Phe Ser Asp Ser Asp Leu Gln
 290 295
Ser Phe Ile Gln Leu Leu Pro Arg Glu Ala His Tyr Thr Lys Lys Phe
              310 315
Leu Gly Cys Val Ala Gln Ser Cys Arg Leu Gly Ile Leu Leu Glu Gly
               325
<210> 573
<211> 426
<212> PRT
<213> C. Trachomatis D serovar
<400> 573
Met Tvr Val Arg Ser Ile Phe Phe Ser Ile Ile Ala Phe Leu Thr Val
1 5
                          10
Gly Cys Ser Phe Ser Pro Pro Glu Ser Gly Leu Ile Ile Ala Ile His
                             25
Asp Asp Pro Arg Ser Leu Ser Pro Glu Lys Gly Glu Asn Ala Phe His
                         40
Phe Ser Leu Ser Lys Ala Leu Phe Ala Thr Leu Phe Arg Glu Glu Leu
                      55
Ser Gly Leu Thr Pro Ala Leu Val Ser Ser Tyr Gln Val Ser Glu Asp
                 70
Gly Arg Phe Tyr Arg Phe Cys Ile Arg Lys Asp Ala Lys Trp Ser Asp
                                 90
Gly Ser Leu Leu Leu Ala Glu Asp Val Ile Ala Ala Trp Glu His Thr
100 105 110
Lys Gln Ala Gly Arg Tyr Ser Leu Leu Phe Glu Lys Leu Ser Phe Arg
115 120 125
Ala Ser Ser Ser Ser Glu Ile Leu Ile Glu Leu Lys Glu Pro Glu Pro
130 135 140
Gln Leu Leu Ala Ile Leu Ala Ser Pro Phe Phe Ala Val Tyr Arg Pro
145 150 155 160
Glu Asn Pro Phe Leu Ser Ser Gly Pro Phe Met Pro Lys Thr Tyr Val
            165 170 175
Gln Gly Gln Thr Leu Val Leu Gln Lys Asn Pro Tyr Tyr Tyr Asp His
                      185
                                                190
```

Ala His Val Glu Leu His Ser Ile Asp Phe Arg Ile Ile Pro Asn Ile

```
200
Tyr Thr Ala Leu His Leu Leu Arg Arg Gly Asp Val Asp Trp Val Gly 210 215 220
Gln Pro Trp His Gln Gly Ile Pro Phe Glu Leu Arg Thr Thr Ser Ala
                  230
                             235
Leu Tyr Thr His Tyr Ser Val Asp Gly Thr Phe Trp Leu Ile Leu Asn 245 250 255
Pro Lys Asp Pro Val Leu Ser Ser Leu Ser Asn Arg Gln Arg Leu Ile
           260 265 270
Ala Ala Val Gln Lys Glu Lys Leu Val Lys Gln Ala Leu Gly Thr Gln
275 280 285
                           280
                                                285
Tyr Arg Val Ala Glu Ser Ser Pro Ser Pro Glu Gly Ile Ile Ala His
                     295
                                            300
Gln Glu Ala Ser Thr Pro Phe Pro Gly Lys Ile Thr Leu Ile Tyr Pro
                    310
                                       315
Asn Asn Ile Thr Arg Cys Gln Arg Leu Ala Glu Val Leu Gln Glu Gln 325 330 335
Cys Arg Asp Ala Gly Ile Gln Leu Thr Leu Glu Gly Leu Glu Tyr His 340 \hspace{1.5cm} 345 \hspace{1.5cm} 350 \hspace{1.5cm}
Val Phe Val Gln Lys Arg Ala Thr Gln Asp Phe Ser Val Ser Thr Ala
355 360 365
Thr Ser Ile Ala Phe His Pro Leu Ala Lys Ser Lys Phe Asp Gln Thr 370 375 380
Ala Leu Asp Asn Phe Thr Cys Leu Pro Leu Tyr His Ile Glu Tyr Asp
                 390
                                      395
Tyr Ile Leu Ser Arg Pro Leu Asp Gln Ile Val His Tyr Pro Ser Gly
            405 410
Ser Val Asp Leu Thr Tyr Ala His Phe His
           420
<210> 574
<211> 605
<212> PRT
<213> C. Trachomatis D serovar
<400> 574
Met Gln Asn Ile Leu Arg Thr Ser Ser Cys Arg Tyr Met Phe Leu Leu
1 5 10 15
Gly Ile Arg Ser Val Trp Asn Arg Val Ala Val Val Asn Asn Phe Arg
Gly Ser Ser Trp Lys Ile Val Ala Ile Pro Ser Cys Ile Leu Phe Thr
                           40
Leu Ile Phe His Leu Pro Arg Trp Leu Ile Asp Phe Gly Val Cys Thr
                        55
Asn Leu Ala Cys Ser Leu Ser Ile Ile Phe Trp Val Phe Ser Leu Arg
                   70
                                        75
Ser Ser Ala Ser Ala Arg Ile Phe Pro Ser Leu Leu Leu Tyr Leu Cys
                                   90
Leu Leu Arg Leu Gly Leu Asn Leu Ala Ser Thr Arg Trp Ile Leu Ser 100 105 110
Ser Gly Trp Ala Ser Pro Leu Ile Phe Ala Leu Gly Asn Phe Phe Ser
115 120 125
Leu Gly Ser Ile Pro Val Ala Leu Thr Val Cys Leu Leu Leu Phe Leu
130 135 140
 130
                                           140
Val Asn Phe Leu Val Ile Thr Lys Gly Ala Glu Arg Ile Ala Glu Val
                 150
                                      155
Arg Ala Arg Phe Ser Leu Glu Ala Leu Pro Gly Lys Gln Met Ser Leu
165 170 175
Asp Ala Asp Ile Ala Ala Gly Arg Ile Gly Tyr Ser Arg Ala Ser Val
           180
                                185
                                                    190
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Lys Lys Ser Ser Leu Leu Glu Glu Ser Asp Tyr Phe Ser Ala Met Glu

200 Gly Val Phe Arg Phe Val Lys Gly Asp Ala Ile Met Ser Trp Val Leu 210 $$ 215 $$ 220 Leu Gly Val Asn Ile Leu Ala Ala Leu Phe Leu Gly Arg Ala Thr His 225 230235235 Val Gly Asp Leu Trp Leu Thr Val Leu Gly Asp Ala Leu Val Ser Gln 245 250 255 Ile Pro Ala Leu Leu Thr Ser Cys Ala Ala Ala Thr Leu Ile Ala Lys 260 265 270 Val Gly Glu Lys Glu Ser Leu Ala Gln His Leu Leu Asp Tyr Tyr Glu 275 280 285 Gln Ser Arg Gln Ser Phe Leu Phe Ile Ala Leu Ile Leu Cys Gly Met 295 300 Ala Cys Ile Pro Gly Ala Pro Lys Ala Leu Ile Leu Gly Phe Ser Val 310 315 Leu Leu Phe Leu Gly Tyr Lys Asn Pro Ser Ser Gly Glu Thr Leu Leu 325 330 335 Phe Gln Lys Glu Arg Val Glu Phe Val Leu Pro Asp Glu Gly Val Gly 340 345 350Asn Pro Ala Asn Leu Tyr Lys Asp Ala Arg Asn Gln Ile Tyr Gln Glu 355 360 365 Leu Gly Val Val Phe Pro Glu Ala Ile Val Val Arg His Val Thr Gly 370 375 380 Ser Ser Pro Arg Leu Ile Phe Ser Gly Gln Glu Val Ala Leu Arg Glu 390 395 Leu Ser Cys Pro Ala Ile Leu Glu Ser Ile Arg Gln Leu Ala Pro Glu
405 410 415 Thr Ile Ser Glu Arg Phe Val Thr Arg Leu Val Asp Glu Phe Arg Glu 420 425 His Ala Phe Leu Ser Ile Glu Glu Ile Leu Pro Leu Lys Ile Ser Glu 435 440 Asn Ser Leu Ile Phe Leu Leu Arg Ala Leu Val Arg Glu Arg Val Ser 450 455 Leu His Leu Phe Pro Lys Ile Leu Glu Ala Ile Asp Val Tyr Gly Ser 465 470 475 480 Gln Pro Lys Asn Ser Gln Glu Leu Val Glu Cys Val Arg Lys Tyr Leu 485 490 495 Gly Lys Gln Ile Gly Leu Ser Leu Trp Asn Arg Gln Asp Val Leu Glu 500 505 510Val Ile Thr Ile Asp Ser Leu Val Glu Gln Phe Val Arg Asp Ser Gln 520 Glu Lys Val Val Leu Asp Leu Asn Glu Lys Val Val Ala Gln Val Lys 535 540 His Leu Leu Arg Val Gly Glu Gly Asn Phe Arg Ala Ile Val Thr Gly 545 550 555 560Ser Glu Thr Arg Lys Glu Leu Lys Arg Ile Val Asp Pro Tyr Phe Pro 565 570 575Asp Leu Leu Val Leu Ala His Ser Glu Leu Pro Glu Glu Ile Pro Ile 580 585 590 Thr Leu Leu Gly Ala Val Ser Asp Glu Val Leu Leu Ser

<210> 575 <211> 173

<400> 575

Met Lys Lys Phe Leu Leu Leu Ser Leu Met Ser Leu Ser Ser Leu Pro 1 5 10 15 15 Thr Phe Ala Ala Asn Ser Thr Gly Thr Ile Gly Ile Val Asn Leu Arg

<212> PRT <213> C. Trachomatis D serovar

<210> 576

<211> 354 <212> PRT

<213> C. Trachomatis D serovar

<400> 576

Met Ser Gln Ser Thr Tyr Ser Leu Glu Gln Leu Ala Asp Phe Leu Lys 10 Val Glu Phe Gln Gly Asn Gly Ala Thr Leu Leu Ser Gly Val Glu Glu Ile Glu Glu Ala Lys Thr Ala His Ile Thr Phe Leu Asp Asn Glu Lys Tyr Ala Lys His Leu Lys Ser Ser Glu Ala Gly Ala Ile Ile Ile Ser Arg Thr Gln Phe Gln Lys Tyr Arg Asp Leu Asn Lys Asn Phe Leu Ile 65 70 75 80Thr Ser Glu Ser Pro Ser Leu Val Phe Gln Lys Cys Leu Glu Leu Phe 85 90 95 Ile Thr Pro Val Asp Ser Gly Phe Pro Gly Ile His Pro Thr Ala Val Ile His Pro Thr Ala Ile Ile Glu Asp His Val Cys Ile Glu Pro Tyr 115 120 Ala Val Val Cys Gln His Ala His Val Gly Ser Ala Cys His Ile Gly 135 140 Ser Gly Ser Val Ile Gly Ala Tyr Ser Thr Val Gly Glu His Ser Tyr 145 150 150 155 160The His Pro Arg Val Val Ile Arg Glu Arg Val Ser Ile Gly Lys Arg 165 170 175 Val Ile Ile Gln Pro Gly Ala Val Ile Gly Ser Cys Gly Phe Gly Tyr 180 185 190 Val Thr Ser Ala Phe Gly Gln His Lys His Leu Lys His Leu Gly Lys 195 200 205 Val Ile Ile Glu Asp Asp Val Glu Ile Gly Ala Asn Thr Thr Ile Asp 210 215 Arg Gly Arg Phe Lys His Ser Val Val Arg Glu Gly Ser Lys Ile Asp 225 230 235 240 Asn Leu Val Gln Ile Ala His Gln Val Glu Val Gly Gln His Ser Met 245 250 255 Ile Val Ala Gln Ala Gly Ile Ala Gly Ser Thr Lys Ile Gly Asn His 260 265 270

Val Ile Ile Gly Gly Gln Ala Gly Ile Thr Gly His Ile Cys Ile Ala

280 Asp His Val Ile Met Met Ala Gln Thr Gly Val Thr Lys Ser Ile Thr 290 295 300 Ser Pro Gly Ile Tyr Gly Gly Ala Pro Ala Arg Pro Tyr Gln Glu Ile 305 310 315 320 His Arg Gln Val Ala Lys Val Arg Asn Leu Pro Arg Leu Glu Glu Arg 325 330 335 Ile Ala Ala Leu Glu Lys Leu Val Gln Lys Leu Glu Ala Leu Ser Glu 345 Gln His <210> 577 <211> 421 <212> PRT <213> C. Trachomatis D serovar Met Thr Ala Ser Gly Gly Ala Gly Gly Leu Gly Ser Thr Gln Thr Val 10 Asp Val Ala Arg Ala Gln Ala Ala Ala Ala Thr Gln Asp Ala Gln Glu 20 25 30 Val Ile Gly Ser Gln Glu Ala Ser Glu Ala Ser Met Leu Lys Gly Cys 35 40 45 Glu Asp Leu Ile Asn Pro Ala Ala Ala Thr Arg Ile Lys Lys Lys Gly 55 Glu Lys Phe Glu Ser Leu Glu Ala Arg Arg Lys Pro Thr Ala Asp Lys Ala Glu Lys Lys Ser Glu Ser Thr Glu Glu Lys Gly Asp Thr Pro Leu Glu Asp Arg Phe Thr Glu Asp Leu Ser Glu Val Ser Gly Glu Asp Phe 105 100 Arg Gly Leu Lys Asn Ser Phe Asp Asp Asp Ser Ser Pro Asp Glu Ile 115 120 125 Leu Asp Ala Leu Thr Ser Lys Phe Ser Asp Pro Thr Ile Lys Asp Leu 130 135 140 Ala Leu Asp Tyr Leu Ile Gln Thr Ala Pro Ser Asp Gly Lys Leu Lys 145 150 155 160 Ser Thr Leu Ile Gln Ala Lys His Gln Leu Met Ser Gln Asn Pro Gln 165 170 175Ala Ile Val Gly Gly Arg Asn Val Leu Leu Ala Ser Glu Thr Phe Ala 180 185 190 Ser Arg Ala Asn Thr Ser Pro Ser Ser Leu Arg Ser Leu Tyr Phe Gln 195 200 Val Thr Ser Ser Pro Ser Asn Cys Ala Asn Leu His Gln Met Leu Ala 210 215 220 Ser Tyr Leu Pro Ser Glu Lys Thr Ala Val Met Glu Phe Leu Val Asn 225 230 235 240 Gly Met Val Ala Asp Leu Lys Ser Glu Gly Pro Ser Ile Pro Pro Ala 245 250 Lys Leu Gln Val Tyr Met Thr Glu Leu Ser Asn Leu Gln Ala Leu His 260 265 270 Ser Val Asn Ser Phe Phe Asp Arg Asn Ile Gly Asn Leu Glu Asn Ser 275 280 285 Leu Lys His Glu Gly His Ala Pro Ile Pro Ser Leu Thr Thr Gly Asn 290 295 300 Leu Thr Lys Thr Phe Leu Gln Leu Val Glu Asp Lys Phe Pro Ser Ser 305 310 315 320Ser Lys Ala Gln Lys Ala Leu Asn Glu Leu Val Gly Pro Asp Thr Gly

330 Pro Gln Thr Glu Val Leu Asn Leu Phe Phe Arg Ala Leu Asn Gly Cys

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340
Ser Pro Arg Ile Phe Ser Gly Ala Glu Lys Lys Gln Gln Leu Ala Ser
                          360
                                          365
Val Ile Thr Asn Thr Leu Asp Ala Ile Asn Ala Asp Asn Glu Asp Tyr
                      375
                                         380
Pro Lys Pro Gly Asp Phe Pro Arg Ser Ser Phe Ser Ser Thr Pro Pro
                  390
                                   395
His Ala Pro Val Pro Gln Ser Glu Ile Pro Thr Ser Pro Thr Ser Thr
              405
                                  410
Gln Pro Pro Ser Pro
           420
<210> 578
<211> 231
<212> PRT
<213> C. Trachomatis D serovar
Met Met Glu Val Phe Met Asn Phe Leu Asp Gln Leu Asp Leu Ile Ile
                                 10
Gln Asn Lys His Met Leu Glu His Thr Phe Tyr Val Lys Trp Ser Lys
         20
                             25
Gly Glu Leu Thr Lys Glu Gln Leu Gln Ala Tyr Ala Lys Asp Tyr Tyr-
                          40
Leu His Ile Lys Ala Phe Pro Lys Tyr Leu Ser Ala Ile His Ser Arg
                      55
 50
Cys Asp Asp Leu Glu Ala Arg Lys Leu Leu Leu Asp Asn Leu Met Asp
                  70
                                      75
Glu Glu Asn Gly Tyr Pro Asn His Ile Asp Leu Trp Lys Gln Phe Val
               85
                                  90
Phe Ala Leu Gly Val Thr Pro Glu Glu Leu Glu Ala His Glu Pro Ser
           100
                              105
Glu Ala Ala Lys Ala Lys Val Ala Thr Phe Met Arg Trp Cys Thr Gly
                          120
Asp Ser Leu Ala Ala Gly Val Ala Ala Leu Tyr Ser Tyr Glu Ser Gln
                      135
Ile Pro Arg Ile Ala Arg Glu Lys Ile Arg Gly Leu Thr Glu Tyr Phe
                                     155
                  150
Gly Phe Ser Asn Pro Glu Asp Tyr Ala Tyr Phe Thr Glu His Glu Glu
                                  170
Ala Asp Val Arg His Ala Arg Glu Glu Lys Ala Leu Ile Glu Met Leu
         180
                              185
Leu Lys Asp Asp Ala Asp Lys Val Leu Glu Ala Ser Gln Glu Val Thr
       195
                        200
Gln Ser Leu Tyr Gly Phe Leu Asp Ser Phe Leu Asp Pro Gly Thr Cys
 210
                    215
Cys Ser Cys His Gln Ser Tyr
<210> 579
<211> 243
<212> PRT
<213> C. Trachomatis D serovar
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<400> 579
Met Lys Ile Thr Pro Ile Lys Thr Arg Lys Val Phe Ala His Asp Ser 1
1 1 5 10
Leu Gln Glu Ile Leu Gln Glu Ala Leu Pro Pro Leu Gln Glu Alg Ser 20
25 30
Val Val Val Val Val Ser Ser Lys Ile Val Ser Leu Cys Glu Gly Ala Val 35
45

Ala Asp Ala Arg Met Cys Lys Ala Glu Leu Ile Lys Lys Glu Ala Asp 50 60 55 Ala Tyr Leu Phe Cys Glu Lys Ser Gly Ile Tyr Leu Thr Lys Lys Glu 65 70 75 80 Gly Ile Leu Ile Pro Ser Ala Gly Ile Asp Glu Ser Asn Thr Asp Gln 85 90 95 Pro Phe Val Leu Tyr Pro Lys Asp Ile Leu Gly Ser Cys Asn Arg Ile 100 105 110 Gly Glu Trp Leu Arg Asn Tyr Phe Arg Val Lys Glu Leu Gly Val Ile 115 120 125 Ile Thr Asp Ser His Thr Thr Pro Met Arg Arg Gly Val Leu Gly Ile 135 140 Gly Leu Cys Trp Tyr Gly Phe Ser Pro Leu His Asn Tyr Ile Gly Ser 150 155 Leu Asp Cys Phe Gly Arg Pro Leu Gln Met Thr Gln Ser Asn Leu Val 165 170 175 Asp Ala Leu Ala Val Ala Ala Val Val Cys Met Gly Glu Gly Asn Glu 180 185 190 Gln Thr Pro Leu Ala Val Ile Glu Gln Ala Pro Asn Met Val Tyr His 195 205 Ser His Pro Thr Ser Arg Glu Glu Tyr Cys Ser Leu Arg Ile Asp Glu 210 215 220 Thr Glu Asp Leu Tyr Gly Pro Phe Leu Gln Ala Val Thr Trp Ser Gln 230 235 Glu Lvs Lvs

<210> 580 <211> 383 <212> PRT

<213> C. Trachomatis D serovar

<400> 580

Met Leu Pro His Gln Gln Asn Ser Ser Ser Glu Arg Ala Arg His His 10 Glu Ser Arg Ser His Arg His Ser Ser Ser Ser Arg His His Val Thr 20 25 Arg Ser Gln Ser Ser Ala Leu Pro Gln Leu Gln Glu Arg Pro Val Pro 40 His Pro Leu Ala Glu Arg Glu Leu Ile Ile Phe His Ser Val His Gln 55 Gln Gln Asn Asn Asn Pro Leu Arg Met Ile Cys Asp Thr Ile Arg Gln 65 70 75 80Ala Gln Arg Gly Ile Phe Met Arg Ile Tyr Thr Ile Ser Ser Asp Asp 85 90 95 Ile Ile Gln Ser Leu Ile Gln Thr Ser His His Val Pro Val Glu Val 105 100 Lys Tyr His Cys Gly Glu Ser Leu Pro Val Ala Cys Gln Asn Ser Arg 115 120 125 Val Val Leu Arg Leu Thr Asn Gly Arg Thr Leu Gln His Lys Lys Thr 135 140 Met Leu Ala Asp Fhe Gln Thr Val Val Thr Gly Ser Ala Asn Tyr Thr 145 150 150 155 160Asp Leu Ser Leu Asn His Asp Ala Asn Val Thr Ala Cys Ile Glu Ser 165 170 Ser Glu Leu His Asp Ala Val Phe Ser Glu Arg Pro Gln Leu Val His 180 185 190 Val Gly Pro Gln Leu Leu Asn Tyr Ile Pro Ile Gln Arg Leu Ile Pro 195 200 205 Asn Ala Ala Ser Lys Met Ile Leu Asn Ala Ile Asn Gln Ala Thr Asp 215 210

PCT/US01/23121

356

Ser Ile Phe Val Leu Met Tyr Ile Phe Leu Ser Pro Glu Phe Phe Leu 235 Ala Leu Ala Gln Ala Met Arg Arg Gly Val Arg Val Lys Val Ile Ile 245 250 Asp Asn His Ser Lys Gln Asp Thr Cys Lys Leu Leu Ser Lys Leu Gly 265 270 Ile Gln Leu Pro Ile Tyr Glu Arg Lys Thr Glu Gly Val Leu His Thr 280 285 Lys Ile Cys Cys Ile Asp Asn Lys Thr Leu Ile Phe Gly Ser Ala Asn 295 300 Trp Ser Gly Ala Gly Met Ile Lys Asn Phe Glu Asp Leu Phe Ile Leu 305 310 315 320 310 315 Arg Pro Ile Thr Glu Thr Gln Leu Gln Ala Phe Met Asp Val Trp Ser 325 330 335 Leu Leu Glu Thr Asn Ser Ser Tyr Leu Ser Pro Glu Ser Val Leu Thr 340 345 350 Ala Pro Thr Pro Ser Ser Arg Pro Thr Gln Gln Asp Thr Asp Ser Asp 360 Asp Glu Gln Pro Ser Thr Ser Gln Gln Asp Ile Arg Met Arg Lys 375 <210> 581 <211> 193 <212> PRT <213> C. Trachomatis D serovar <400> 581 Met Trp Phe Phe Leu Gly Ser Pro Ser Ala Ile Thr Asn Phe Ser Arg 10 20 25

Val Asp Val Ala Leu Asn Leu Arg Ile Asn Arg Gln Ile Arg Ala Pro Arg Val Arg Val Ile Gly Ser Ala Gly Glu Gln Leu Gly Ile Leu Ser 40 Ile Lys Glu Ala Leu Asp Leu Ala Lys Glu Ala Asn Leu Asp Leu Val 55 Glu Val Ala Ser Asn Ser Glu Pro Pro Val Cys Lys Ile Met Asp Tyr 70 75 Gly Lys Tyr Arg Tyr Asp Val Thr Lys Lys Glu Lys Asp Ser Lys Lys 85 90 95 Ala Gln His Gln Val Arg Ile Lys Glu Val Lys Leu Lys Pro Asn Ile 100 105 Asp Asp Asn Asp Phe Leu Thr Lys Ala Lys Gln Ala Arg Ala Phe Ile 120 125 Glu Lys Gly Asn Lys Val Lys Val Ser Cys Met Phe Arg Gly Arg Glu 135 140 Leu Ala Tyr Pro Glu His Gly Tyr Lys Val Ile Gln Arg Met Cys Gln 150 155 Gly Leu Glu Asp Ile Gly Phe Val Glu Ser Glu Pro Lys Leu Asn Gly 165 170 Arg Ser Leu Ile Cys Val Ile Ala Pro Gly Thr Leu Lys Thr Lys Lys 180 185 Lys

<210> 582

<211> 264 <212> PRT

<213> C. Trachomatis D serovar

<400> 582

Met Gly Asn Ser Gly Phe Tyr Leu Tyr Asn Thr Glu Asn Cys Val Phe

10 Ala Asp Asn Ile Lys Val Gly Gln Met Thr Glu Pro Leu Lys Asp Gln 20 25 30 Gln Ile Ile Leu Gly Thr Lys Ser Thr Pro Val Ala Ala Lys Met Thr 40 Ala Ser Asp Gly Ile Ser Leu Thr Val Ser Asn Asn Ser Ser Thr Asn 55 Ala Ser Ile Thr Ile Gly Leu Asp Ala Glu Lys Ala Tyr Gln Leu Ile 75 Leu Glu Lys Leu Gly Asn Gln Ile Leu Asp Gly Ile Ala Asp Thr Ile 8.5 90 Val Asp Ser Thr Val Gln Asp Ile Leu Asp Lys Ile Thr Thr Asp Pro 100 105 Ser Leu Gly Leu Leu Lys Ala Phe Asn Asn Phe Pro Ile Thr Asn Lys 120 Ile Gln Cys Asn Gly Leu Phe Thr Pro Ser Asn Ile Glu Thr Leu Leu 135 Gly Gly Thr Glu Ile Gly Lys Phe Thr Val Thr Pro Lys Ser Ser Gly 145 150 155 160 Ser Met Phe Leu Val Ser Ala Asp Ile Ile Ala Ser Arg Met Glu Gly 165 170 175 Gly Val Val Leu Ala Leu Val Arg Glu Gly Asp Ser Lys Pro Cys Ala 180 185 190 Ile Ser Tyr Gly Tyr Ser Ser Gly Val Pro Asn Leu Cys Ser Leu Arg Thr Ser Ile Thr Asn Thr Gly Leu Thr Pro Thr Thr Tyr Ser Leu Arg 210 215 220 Val Gly Gly Leu Glu Ser Gly Val Val Trp Val Asn Ala Leu Ser Asn 225 230 235 240 Gly Asn Asp Ile Leu Gly Ile Thr Asn Thr Ser Asn Val Ser Phe Leu 245 250 255 250 245 Glu Val Ile Pro Gln Thr Asn Ala

Met Phe Thr Arg Ile Val Met Val Asp Leu Gln Glu Lys Gln Cys Thr 10 Ile Val Lys Arg Asn Gly Met Phe Val Pro Phe Asp Arg Asn Arg Ile 25 Phe Gln Ala Leu Glu Ala Ala Phe Arg Asp Thr Arg Arg Ile Asp Asp 40 His Met Pro Leu Pro Glu Asp Leu Glu Ser Ser Ile Arg Ser Ile Thr 55 His Gln Val Val Lys Glu Val Val Gln Lys Ile Thr Asp Gly Gln Val 65 70 70 75 80Val Thr Val Glu Arg Ile Gln Asp Met Val Glu Ser Gln Leu Tyr Val 85 90 95 Asn Gly Leu Gln Asp Val Ala Arg Asp Tyr Ile Val Tyr Arg Asp Asp 100 105 110 105 Arg Lys Ala His Arg Lys Lys Ser Trp Gln Ser Leu Ser Val Val Arg 120 Arg Cys Gly Thr Val Val His Phe Asn Pro Met Lys Ile Ser Ala Ala 130 135

Leu Glu Lys Ala Phe Arg Ala Thr Asp Lys Thr Glu Gly Met Thr Pro

Ser Ser Val Arg Glu Glu Ile Asn Ala Leu Thr Gln Asn Ile Val Ala

155

<210> 583 <211> 1053

<212> PRT <400> 583

<213> C. Trachomatis D serovar

				165					170					175	
Glu	Ile	Glu	Glu 180	Cys	Cys	Pro	Gln	Gln 185	Asp	Arg	Arg	Ile	Asp 190	Ile	Glu
Lys	Ile	G1n 195	Asp	Ile	Val	Glu	Gln 200	Gln	Leu	Met	Val	Val 205	Gly	His	Tyr
Ala	Val 210	Ala	Lys	Asn	Tyr	Ile 215	Leu	Tyr	Arg	Glu	Ala 220	Arg	Ala	Arg	Val
Arg 225	Asp	Asn	Arg	Glu	Glu 230	Asp	Gly	Ser	Thr	Glu 235	Lys	Thr	Ile	Ala	Glu 240
Glu	Ala	Val	Glu	Val 245	Leu	Ser	Lys	Asp	Gly 250	Ser	Thr	Tyr	Thr	Met 255	Thr
His	Ser	Gln	Leu 260	Leu	Ala	His	Leu	Ala 265	Arg	Ala	Суѕ	Ser	Arg 270	Phe	Pro
		275					280					285		Asn	
	290			-		295					300	-		Met	
305					310				-	315				Ala	320
				325			-	-	330			-	-	Ser 335	-
			340					345					350	Arg	
		355					360					365		His	
	370		-			375	_			_	380		-	Asp	
385	Phe				390					395				Phe	400
				405					410					Met 415	
			420				Asn	425					430		
Ile		435					440					445		Ala	
Pro	450					455		-			460			Ser	-
465	Leu				470					475		-	-	Val	480
Ala				485					490					Asn 495	
Trp			500					505			-	_	510	Asn	_
-		515	-				520		-			525	-	Thr	
	530					535	Lys				540				Tyr
545	Glu		•		550					555				Arg	560
			_	565					570	-				Ala 575	
Trp			580					585				_	590	Thr	
		595					600					605		Tyr	
	610					615					620	-		Asp -	
625					630					635		-		Trp	640
Lys	Met	Leu	Ser	Met 645	Leu	Phe	Glu	Thr	Gly 650	His	Pro	Trp	Met	Thr 655	Phe

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Lys Asp Pro Ser Asn Ile Arg Ser Ala Gln Asp His Lys Gly Val Val
660 665 670
Arg Cys Ser Asn Leu Cys Thr Glu Ile Leu Leu Asn Cys Ser Glu Thr
              680
Glu Thr Ala Val Cys Asn Leu Gly Ser Ile Asn Leu Val Gln His Ile
690 695 700
Val Gly Asp Gly Leu Asp Glu Glu Lys Leu Ser Glu Thr Ile Ser Ile
705 710 715 720
Ala Val Arg Met Leu Asp Asn Val Ile Asp Ile Asn Phe Tyr Pro Thr
725 730 735
Lys Glu Ala Lys Glu Ala Asn Phe Ala His Arg Ala Ile Gly Leu Gly 740 745 750
Val Met Gly Phe Gln Asp Ala Leu Tyr Lys Leu Asp Ile Ser Tyr Ala
                             760
        755
                                                   765
Ser Gln Glu Ala Val Glu Phe Ala Asp Tyr Ser Ser Glu Leu Ile Ser
                          775
                                                780
Tyr Tyr Ala Ile Gln Ala Ser Cys Leu Leu Ala Lys Glu Arg Gly Thr
                                       795
                  790
Tyr Ser Ser Tyr Lys Gly Ser Lys Trp Asp Arg Gly Leu Leu Pro Ile
805 810 815
Asp Thr Ile Gln Leu Leu Ala Asn Tyr Arg Gly Glu Ala Asn Leu Gln
820 825 830
Met Asp Thr Ser Ser Arg Lys Asp Trp Glu Pro Ile Arg Ser Leu Val
835 840 845
Lys Glu His Gly Met Arg His Cys Gln Leu Met Ala Ile Ala Pro Thr
                         855
                                               860
Ala Thr Ile Ser Asn Ile Ile Gly Val Thr Gln Ser Ile Glu Pro Thr
                    870
                                         875
Tyr Lys His Leu Phe Val Lys Ser Asn Leu Ser Gly Glu Phe Thr Ile
885 890 895
Pro Asn Val Tyr Leu Ile Glu Lys Leu Lys Lys Leu Gly Ile Trp Asp
900 905 910
Ala Asp Met Leu Asp Asp Leu Lys Tyr Phe Asp Gly Ser Leu Leu Glu 915 920 925
Ile Glu Arg Ile Pro Asp His Leu Lys His Ile Phe Leu Thr Ala Phe
                         935
Glu Ile Glu Pro Glu Trp Île Ile Glu Cys Ala Ser Arg Arg Gln Lys
945 950 955 960
Trp Ile Asp Met Gly Gln Ser Leu Asn Leu Tyr Leu Ala Gln Pro Asp
965 970 975
Gly Lys Lys Leu Ser Asn Met Tyr Leu Thr Ala Trp Lys Lys Gly Leu
980 985 990
Lys Thr Thr Tyr Tyr Leu Arg Ser Ser Ser Ala Thr Thr Val Glu Lys
995 1000 1005
Ser Phe Val Asp Ile Asn Lys Arg Gly Ile Gln Pro Arg Trp Met Lys 1010 1015 1020
Asn Lys Ser Ala Ser Ala Gly Ile Ile Val Glu Arg Ala Lys Lys Ala
1025 1030 1035 1040
Pro Val Cys Ser Leu Glu Glu Gly Cys Glu Ala Cys Gln
                1045
                                      1050
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<210> 584 <211> 346

<400> 584

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<211> 346 <212> PRT

<213> C. Trachomatis D serovar

Pro Ile Lys Tyr Lys Trp Ala Trp Glu His Tyr Leu Asn Gly Cys Ala 35 40 45 Asn Asn Trp Leu Pro Thr Glu Ile Pro Met Gly Lys Asp Ile Glu Leu 55 Trp Lys Ser Asp Arg Leu Ser Glu Asp Glu Arg Arg Val Ile Leu Leu 65 70 75 80 75 Asn Leu Gly Phe Phe Ser Thr Ala Glu Ser Leu Val Gly Asn Asn Ile Val Leu Ala Ile Phe Lys His Val Thr Asn Pro Glu Ala Arg Gln Tyr 100 ' 105 Leu Leu Arg Gln Ala Phe Glu Glu Ala Val His Thr His Thr Phe Leu 120 125 115 Tyr Ile Cys Glu Ser Leu Gly Leu Asp Glu Lys Glu Ile Phe Asn Ala 130 135 140 Tyr Asn Glu Arg Ala Ala Ile Lys Ala Lys Asp Asp Phe Gln Met Glu 145 150 155 160 Ile Thr Gly Lys Val Leu Asp Pro Asn Phe Arg Thr Asp Ser Val Glu 165 · 170 175 Gly Leu Gln Glu Phe Val Lys Asn Leu Val Gly Tyr Tyr Ile Ile Met 180 185 190 Glu Gly Ile Phe Phe Tyr Ser Gly Phe Val Met Ile Leu Ser Phe His 195 200 205 Arg Gln Asn Lys Met Ile Gly Ile Gly Glu Gln Tyr Gln Tyr Ile Leu 210 215 220 Arg Asp Glu Thr Ile His Leu Asn Phe Gly Ile Asp Leu Ile Asn Gly 225 230 235 240 235 Ile Lys Glu Glu Asn Pro Glu Ile Trp Thr Pro Glu Leu Gln Glu Glu 245 250 255Ile Val Glu Leu Ile Lys Arg Ala Val Asp Leu Glu Ile Glu Tyr Ala 265 Gln Asp Cys Leu Pro Arg Gly Ile Leu Gly Leu Arg Ala Ser Met Phe 275 280 285 Ile Asp Tyr Val Gln His Ile Ala Asp Arg Arg Leu Glu Arg Ile Gly 290 295 300 Leu Lys Pro Ile Tyr His Thr Lys Asn Pro Phe Pro Trp Met Ser Glu 305 310 315 320 Thr Ile Asp Leu Asn Lys Glu Lys Asn Phe Phe Glu Thr Arg Val Ile 325 330 335 Glu Tyr Gln His Ala Ala Ser Leu Thr Trp 340

<400> 585

X-40U5 585 Met Ser Phe Phe His Thr Arg Lys Tyr Lys Leu Ile Leu Arg Gly Leu Il Leu Cys Leu Ala Gly Cys Phe Leu Met Asn Ser Cys Ser Ser Ser Arg Gly Asn Gln Pro Ala Asp Glu Ser Ile Tyr Val Leu Ser Met Asn Arg 35 Met Ile Cys Asp Cys Val Ser Arg Ile Thr Gly Asp Arg Val Lys Asn 50 Ser Val Leu Ile Asp Gly Ala Ile Asp Pro His Ser Tyr Glu Met Val Cys Asp Cys Val Ser Arg Ile Thr Gly Asp Arg Val Lys Asn 50 Lys Gly Asp Glu Asp Arg Met Ala Met Ser Gln Leu Ile Phe Cys Asn 85 Gly Asp Cys Asp Cys Asp Cys Asp Cys Gly Leu Gly Leu Glu His Ser Ala Ser Leu Arg Lys His Leu Glu Gly Leu Gly Leu Gly Leu Glu His Ser Ala Ser Leu Arg Lys His Leu Glu Gly 105

<210> 585 <211> 326

<212> PRT

<213> C. Trachomatis D serovar

Asn Pro Lys Val Val Asp Leu Gly Gln Arg Leu Leu Asn Lys Asn Cys 115 120 125Phe Asp Leu Leu Ser Glu Glu Gly Phe Pro Asp Pro His Ile Trp Thr 135 140 Asp Met Arg Val Trp Gly Ala Ala Val Lys Glu Met Ala Ala Ala Leu 145 150 155 160 155 Ile Gln Gln Phe Pro Gln Tyr Glu Glu Asp Phe Gln Lys Asn Ala Asp 170 175 165 Leu Ser Thr Ile Pro Glu Lys Asn Arg Tyr Leu Val Thr Gly His Asn 195 200 205 Ala Phe Ser Tyr Phe Thr Arg Arg Tyr Leu Ser Ser Asp Ala Glu Arg 210 215 220 Val Ser Gly Glu Trp Arg Ser Arg Cys Ile Ser Pro Glu Gly Leu Ser 225 230 240 Pro Glu Ala Gln Ile Ser Ile Arg Asp Ile Met Arg Val Val Glu Tyr 245 250 Ile Ser Ala Asn Asp Val Glu Val Val Phe Leu Glu Asp Thr Leu Asn 260 265 270 Gln Asp Ala Leu Arg Lys Ile Val Ser Cys Ser Lys Ser Gly Gln Lys 275 280 285 Ile Arg Leu Ala Lys Ser Pro Leu Tyr Ser Asp Asn Val Cys Asp Asn 290 295 300 Tyr Phe Ser Thr Phe Gln His Asn Val Arg Thr Ile Thr Glu Glu Leu 305 310 315 320 Gly Gly Thr Val Leu Glu

<400> 586

Val Gly Cys Asn Leu Ala Gln Phe Leu Gly Lys Lys Val Leu Leu Ala 1 1 5 10 15 Asp Leu Asp Pro Gln Ser Asn Leu Ser Ser Gly Leu Gly Ala Ser Val 20 25 30 Arg Asn Asn Gln Lys Gly Leu His Asp Ile Val Tyr Lys Ser Asn Asp

<210> 586

<211> 102

<212> PRT

<213> C. Trachomatis D serovar

<210> 587

<211> 243

<212> PRT

<213> C. Trachomatis D serovar

<400> 587

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<210> 588 <211> 527 <212> PRT

<213> C. Trachomatis D serovar

<400> 588 Met Pro Ser Leu Ser Gln Ser Arg Arg Ile Ile Gln Gln Ser Ser Ile

10 Arg Lys Ile Trp Asn Gln Ile Asp Thr Ser Pro Lys His Gly Val Cys 25 30Val Pro Leu Phe Ser Leu Tyr Thr Gln Glu Ser Cys Gly Ile Gly Glu 35 40 45 Phe Leu Asp Leu Ile Pro Met Ile Asp Trp Cys Ile Ser Cys Gly Phe 55 60 Gln Ile Leu Gln Ile Leu Pro Ile Asn Asp Thr Gly Ser Cys Ser Ser 70 75 Pro Tyr Asn Ser Ile Ser Ser Ile Ala Leu Asn Pro Leu His Leu Ser 85 90 Ile Ser Ala Leu Pro Tyr Lys Glu Glu Val Pro Ala Ala Glu Thr Arg Ile Arg Glu Met Gln Gln Leu Ser Gln Leu Pro Gln Val His Tyr Glu 115 120 125 Lys Val Arg Ser Met Lys Arg Asp Phe Phe Gln Glu Tyr Tyr Arg Val 130 135 140 Cys Lys Gln Lys Lys Leu Thr Asp His Pro Asp Phe Tyr Ala Phe Cys 145 150 155 160 Glu Gln Glu Lys Tyr Trp Leu His Pro Tyr Ala Leu Phe Arg Ser Ile 165 170 175 Arg Glu His Leu Asp Asn Leu Pro Ile Asn His Trp Pro Thr Thr Tyr 180 185 190 Thr Asp Leu Ser Gln Ile Thr Glu His Glu Arg Thr Phe Ala Glu Asp 200

Ile Gln Phe His Ser Tyr Leu Gln Tyr Leu Cys Phe Gln Gln Met Thr

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215
                                             220
Gln Val Arg Glu His Ala Asn Cys Lys Ser Cys Leu Ile Lys Gly Asp
225 230 235 240
             230
Ile Fro Ile Leu Ile Ser Lys Asp Ser Cys Asp Val Trp Phe Tyr Arg
             245
                                250
His Tyr Phe Ser Ser Ser Glu Ser Val Gly Ala Pro Pro Asp Leu Tyr 260 265 270
Asn Ala Glu Gly Gln Asn Trp His Leu Pro Ile Cys Asn Met Lys Thr
                280
Leu Gln Gln Asp Asn Tyr Leu Trp Trp Lys Glu Arg Leu Arg Tyr Ala
290 295 300
                      295
Glu Asn Phe Tyr Ser Leu Tyr Arg Leu Asp His Val Val Gly Leu Phe
305 310 315 320
Arg Phe Trp Val Trp Asp Glu Ser Gly Cys Gly Arg Phe Glu Pro His 325 330 335
                                    330
Asp Pro Lys Asn Tyr Leu Ala Gln Gly Gln Asp Ile Leu Ser His Leu
                               345
Leu Thr Ser Ser Ser Met Leu Pro Ile Gly Glu Asp Leu Gly Thr Ile
                            360
Pro Ser Asp Val Lys Arg Met Leu Glu Ser Phe Ala Val Cys Gly Thr
                      375
Arg Ile Pro Arg Trp Glu Arg Asn Trp Glu Gly Asn Gly Ala Tyr Thr
385 390 395 400
                                       395
Pro Phe Asp Gln Tyr Asp Pro Leu Ser Val Thr Ser Leu Ser Thr His 405 410 415
Asp Ser Ser Thr Leu Ala Ser Trp Trp Lys Glu Ser Pro Gln Glu Ser
          420
                               425
Lys Leu Phe Ala Gln Phe Leu Gly Leu Pro Tyr Ser Ser Thr Leu Ser
                           440
Leu His Asn His Thr Glu Ile Leu Lys Leu Ser His Lys Thr Ser Ser 450 455
Ile Phe Arg Ile Asn Leu Ile Asn Asp Tyr Leu Ala Leu Phe Pro Asp
                   470
                                       475
Leu Ile Ser Lys Thr Pro Arg Tyr Glu Arg Ile Asn Leu Pro Gly Thr
               485
                               490
                                                      495
Ile Ser Lys Asn Asn Trp Val Tyr Arg Val Lys Pro Ser Ile Glu Asp
500 505 510
Leu Ser Ser His Ser Lys Leu Asn Ser Leu Leu Glu Ala Leu Phe
<210> 589
<211> 146
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<212> PRT

<213> C. Trachomatis D serovar

<400> 589

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Ser Ile Ser Lys Gly Ala Leu Pro Asp Leu His Ala Leu Gly Met Tyr
His Leu
145
<210> 590
<211> 650
<212> PRT
<213> C. Trachomatis D serovar
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1 10 15
Arg Thr Ile Gly Phe Arg Leu Trp Leu Ile Cys Val Ala Ala Ile Met
Phe Pro Leu Gly Ile Asn Ile Leu Gln Leu Asn Leu Gln Gln Tyr Lys
                                40
Lys Thr Leu Ser Ser Ile Thr Ser Asp Leu Arg Glu Asn Ala Leu Phe 50 55 60
Lys Ala His Thr Leu Gln Gln Thr Ile Pro Leu Asn Ile Asp Ile Leu
65 70 75 80
Ala Leu Phe Ser Glu Ile Phe Asp Leu Asp Arg Gly Val Pro Ala Glu
85 90 95
Pro Asp Leu Ala Leu Ser Lys Glu Met Glu Lys Ile Phe His Ser Thr
100 105 110
Tyr Lys Glu Ile Ser Leu Val Lys Lys Glu Ala Asp Gly Asn Phe Arg
115 120 125
                                120
Val Val Ala Ser Ser Arg Ile Glu Gln Leu Gly Lys Asn Tyr Asn Gln
130 135 140
Glu Ile Phe Leu Ser Asp Ser Gln Pro Phe Leu Ala Thr Leu Arg His
145 150 150 155 160
                                              155
Ser Gly Ser Asp Ser Gln Val Leu Ala Val Leu Gln Thr Asn Ile Phe
165 170 175
Asp Ile Ser Ser Gln Glu Val Leu Gly Val Leu Tyr Thr Leu Ser Asp
180 185 190
Thr Asn Tyr Leu Leu Asn Gly Leu Leu Ala Ala Lys Asp Pro Leu Ser
195 200 205
Val Lys Thr Ala Ile Leu Ser Lys Asn Gly Ile Ile Leu Gln Ala Thr
210 215 220
Asp Ser Ser Leu Asp Leu Val Ser Ile His Lys Thr Val Ser Lys Glu
225 230 235 240
Gln Phe Cys Asp Val Phe Leu Arg Asp Asp Ile Cys Pro Pro His Leu
245 250 255
                  245
Leu Leu Arg Pro Pro Leu Asn Leu Asp Pro Leu Pro Tyr Gly Glu Asn 260 \hspace{1.5cm} 265 \hspace{1.5cm} 265 \hspace{1.5cm} 270 \hspace{1.5cm}
Phe Val Ser Phe Cys Ile Gly Asn Thr Glu Met Trp Gly Tyr Ile His 275 280 285
Ser Leu Pro Glu Met Asp Phe Arg Ile Leu Thr Tyr Glu Glu Lys Ser 290 295 300
Ile Ile Phe Ala Ser Leu Trp Arg Arg Thr Leu Leu Tyr Phe Ala Tyr 305 310 315 320
Phe Cys Cys Val Leu Leu Gly Ser Ile Thr Ala Phe Leu Val Ala Lys
325 330 335
Arg Leu Ser Lys Pro Ile Arg Lys Leu Ala Thr Ala Met Met Glu Thr 340 345 350
Arg Arg Asn Gln His His Pro Tyr Glu Pro Asp Ser Leu Gly Phe Glu 355 360 365
Ile Asn His Leu Gly Glu Ile Phe Asn Ser Met Val Gln Ser Leu Leu 370 375 380
Gln Gln Gln Ser Leu Ala Glu Lys Asn Phe Glu Ile Lys Gln His Ala
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390 Gln Asn Ala Leu Arg Leu Gly Glu Glu Ala Gln Gln Cys Leu Leu Pro 405 410 415Asn Gln Leu Pro Asp Ser Pro Thr Thr Glu Ile Ala Lys Ala Tyr Ile 420 425 430Pro Ala Ile Thr Val Gly Gly Asp Phe Phe Asp Ile Phe Val Ile Gly
435 440 445 Glu Gly Pro Gln Ala Lys Leu Phe Leu Ile Val Ala Asp Ala Ser Gly 450 455 460 Lys Gly Val Asn Ala Cys Ala Tyr Ser Leu Phe Leu Lys Asn Met Leu 465 470 480 His Thr Phe Leu Ser Glu Leu Ser Ser Ile Gln Glu Ala Val Gln Gln 485 490 495 Thr Ala Ala Leu Phe Tyr Gln Gln Thr Ala Glu Ser Gly Met Phe Val 500 505 Thr Leu Cys Ile Tyr Cys Tyr His Tyr Ala Thr Arg Glu Leu Glu Tyr 515 520 525 520 Tyr Ser Cys Gly His Asn Pro Ala Cys Leu Arg Ala Pro Asn Gly Asp 530 535 540 Ile Ser Phe Leu Ser His Pro Gly Met Ala Leu Gly Phe Leu Pro Glu 545 550 560Val Pro Pro His Pro Ala Tyr Thr Leu Val Leu Glu Glu Glu Ser Leu 565 570 575 Leu Val Leu Tyr Thr Asp Gly Val Thr Glu Ala Ser Asn Lys His Gly 580 585 590 Glu Met Phe Gly Glu Glu Arg Leu Lys Ala Leu Val Ala Ser Leu Thr 595 600 605 Lys Gln Ser Ala Glu Glu Ala Ile Gln Ser Ile Met Phe Ser Ile Lys 615 620 Ser Phe Val Lys Asp Cys Pro Gln His Asp Asp Ile Thr Leu Leu Val 625 630 635 Leu Lys Ile Pro Lys Glu Pro Ser Ala Tyr <210> 591 <211> 313 <212> PRT <213> C. Trachomatis D serovar <400> 591 Met Leu Ser Tyr Ile Lys Arg Arg Leu Leu Phe Asn Leu Leu Ser Leu 5 10 Trp Val Val Val Thr Leu Thr Phe Phe Ile Ile Lys Thr Ile Pro Gly 20 25 Asp Pro Phe Asn Asp Glu Asn Gly Asn Ile Leu Ser Ser Glu Thr Leu 35 40 45 Ala Leu Leu Lys Asn Arg Tyr Gly Leu Asp Lys Pro Leu Phe Thr Gln 50 60Tyr Leu Ile Tyr Leu Lys Cys Leu Leu Thr Leu Asp Phe Gly Glu Ser 65 70 75 80 Leu Ile Tyr Lys Asp Arg Thr Val Ile Ser Ile Ile Ala Ala Ala Leu 85 90 Pro Ser Ser Ala Ile Leu Gly Leu Glu Ser Leu Cys Leu Ser Leu Phe $100 \hspace{1cm} 105 \hspace{1cm} 110 \hspace{1cm}$ Gly Gly Ile Thr Leu Gly Ile Leu Ala Ala Phe Tyr Lys Lys Ser Cys 115 120 125 Gly Arg Thr Ile Phe Phe Ser Ser Val Ile Gln Ile Ser Val Pro Ala 130 135 140 Phe Val Ile Gly Ala Phe Leu Gln Tyr Val Phe Ala Ile Lys Tyr Ser 150 155

Cys Leu Pro Ile Ala Cys Trp Gly Asn Phe Ser His Thr Leu Leu Pro

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170

Ser Ile Ala Leu Ala Ile Thr Pro Met Ala Phe Ile Thr Gln Leu Thr 185 180 Cys Ala Ser Val Ser Ala Asn Leu Lys Lys Asp Tyr Val Leu Leu Ala 200 Tyr Ala Lys Gly Leu Ser Pro Phe Lys Val Leu Ile Lys His Ile Leu 210 215 220 Pro Tyr Ala Leu Phe Pro Val Ile Ser Tyr Ser Ala Phe Leu Ile Thr Thr Leu Met Thr Gly Thr Phe Ser Ile Glu Asn Leu Phe Cys Ile Pro 245 250 255 Gly Leu Gly Lys Trp Phe Ile Cys Ser Ile Lys Gln Arg Asp Tyr Pro 260 265 270 Ile Thr Leu Gly Leu Ser Val Phe Tyr Gly Ala Phe Phe Met Leu Thr 275 280 285 Ser Leu Cys Cys Asp Leu Leu Gln Ala Trp Ile Asp Pro Gln Ile Arg 290 295 300

Tyr Ser Tyr Gly Lys Glu Arg Ser Lys 310

<210> 592 <211> 1237 <212> PRT

<213> C. Trachomatis D serovar

Met Thr Trp Ile Pro Leu His Cys His Ser Gln Tyr Ser Ile Leu Asp 10 Ala Thr Cys Ser Ile Lys Lys Phe Val Ala Lys Ala Val Glu Tyr Gln 20 25 Ile Pro Ala Leu Ala Leu Thr Asp His Gly Asn Leu Phe Gly Ala Val 40 Glu Phe Tyr Lys Thr Cys Lys Gln Asn Ala Ile Lys Pro Ile Ile Gly 50 55 60 Cys Glu Leu Tyr Val Ala Pro Ser Ser Arg Phe Asp Lys Lys Lys Glu Arg Lys Ser Arg Val Ala Asn His Leu Ile Leu Leu Cys Lys Asp Glu 85 90 95Glu Gly Tyr Arg Asn Leu Cys Leu Leu Ser Ser Leu Ala Tyr Thr Glu 100 105 110 Gly Phe Tyr Tyr Val Pro Arg Ile Asp Arg Asp Leu Leu Ser Gln His 115 120 125 Ser Lys Gly Leu Ile Cys Leu Ser Ala Cys Leu Ser Gly Ser Val Ala 130 135 140 Gln Ala Ala Leu Glu Ser Glu Glu Asp Leu Glu Lys Asp Leu Leu Trp 145 150 155 160 Tyr Gln Asp Leu Phe Gln Glu Asp Phe Phe Ser Glu Val Gln Leu His 165 170 Lys Ser Ser Glu Glu Lys Val Ala Leu Phe Glu Glu Thr Trp Leu Lys 185 190 Gln Asn Tyr Tyr Gln Phe Ile Glu Lys Gln Leu Lys Val Asn Glu Ala 200 Val Leu Ala Thr Ser Lys Arg Leu Gly Ile Pro Ser Val Ala Thr Asn 210 215 220 Asp Ile His Tyr Leu Asn Pro Asp Asp Trp Leu Ala His Glu Ile Leu 225 230 235 240 Leu Asn Val Gln Ser Arg Glu Pro Ile Arg Thr Ala Lys Gln Asn Thr 245 250 255 Tyr Ile Pro Asn Pro Lys Arg Lys Thr Tyr Pro Ser Arg Glu Phe Tyr 265

Phe Lys Ser Pro Gln Glu Ile Ala Glu Leu Phe Ala Ala His Pro Glu

Thr Ile Thr Asn Thr Cys Ile Val Ala Glu Arg Cys His Leu Glu Leu 290 295 300 Asp Phe Glu Thr Lys His Tyr Pro Ile Tyr Val Pro Glu Ala Leu Gln 310 315 Lys Lys Gly Ser Tyr Thr Glu Glu Glu Arg Tyr Lys Ala Ser Ser Ala 325 330 335 Phe Leu Glu Glu Leu Cys Glu Gln Gly Leu Thr Ser Lys Tyr Thr Pro 340 345 350 Glu Leu Leu Gly His Ile Ala Lys Lys Phe Pro Gly Glu Asp Pro Leu 355 360 365 Thr Leu Val Lys Glu Arg Leu Lys Leu Glu Ser Ser Ile Ile Ile Ser 375 380 Lys Gly Met Cys Asp Tyr Leu Leu Ile Val Trp Asp Ile Ile Asn Trp 385 390 395 400 Ala Lys Asp His Gly Ile Pro Val Gly Pro Gly Arg Gly Ser Gly Ala
405 410 415 Gly Ser Val Met Leu Phe Leu Leu Gly Ile Thr Glu Ile Glu Pro Ile $420 \hspace{1.5cm} 425 \hspace{1.5cm} 430$ Arg Phe Asp Leu Phe Phe Glu Arg Phe Ile Asn Pro Glu Arg Ile Ser 435 440 445 Tyr Pro Asp Ile Asp Ile Asp Ile Cys Met Ile Gly Arg Glu Arg Val 455 460 Ile Asn Tyr Ala Ile Glu Arg His Gly Lys Asp Asn Val Ala Gln Ile Ile Thr Phe Gly Thr Met Lys Ala Lys Met Ala Ile Lys Asp Val Gly
485 490 495 Arg Thr Leu Asp Thr Pro Leu Ala Lys Val Asn Phe Ile Ala Lys His 500 505 510Ile Pro Asp Leu Asn Ala Thr Ile Thr Ser Ala Leu Glu Ala Asp Pro 515 520 525 Glu Leu Arg Gln Leu Tyr Val Asp Asp Ala Glu Ala Ala Glu Val Ile 530 540 Asp Met Ala Lys Lys Leu Glu Gly Ser Ile Arg Asn Thr Gly Val His 545 550 555 560Ala Ala Gly Val Ile Ile Cys Gly Asp Pro Leu Thr Asn His Ile Pro 565 570 575Ile Cys Val Pro Lys Asp Ser Ser Met Ile Ser Thr Gln Tyr Ser Met 580 585 590 Lys Pro Val Glu Ser Val Gly Met Leu Lys Val Asp Phe Leu Gly Leu 595 600 605 Lys Thr Leu Thr Gly Ile His Ile Ala Thr Gln Ala Ile Tyr Lys Lys 615 620 Thr Gly Ile Leu Leu Arg Ala Ala Thr Ile Pro Leu Asp Asp Gln Asn 625 630 635 630 635 Thr Phe Ser Leu Leu His Gln Gly Lys Thr Met Gly Ile Phe Gln Met 645 650 655Glu Ser Arg Gly Met Gln Asp Leu Ala Lys Asn Leu Arg Pro Asp Ala 660 665 670 Phe Glu Glu Ile Ile Ala Ile Gly Ala Leu Tyr Arg Pro Gly Pro Met 675 680 685 Asp Met Ile Pro Ser Phe Ile Asn Arg Lys His Gly Lys Glu Asn Ile 690 695 700 Glu Tyr Asp His Pro Leu Met Glu Pro Ile Leu Lys Glu Thr Phe Gly 705 710 715 720 Ile Met Val Tyr Gln Glu Gln Val Met Gln Ile Ala Gly Ser Leu Ala 725 730 735 Lys Tyr Ser Leu Gly Glu Gly Asp Val Leu Arg Arg Ala Met Gly Lys 740 750 Lys Asp His Glu Gln Met Val Lys Glu Arg Glu Lys Phe Cys Ser Arg 755 760 765

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				_										
Ala Ala 770	A1a	Asn	G1y	Ile	775	Pro	Ser	Ile	Ala	Thr 780	Thr	Ile	Phe	Asp
Lys Met 785	Glu	Lys	Phe	A1a 790	Ser	Tyr	G1y	Phe	Asn 795	Lys	Ser	His	Ala	Ala 800
Ala Tyr	G1y	Leu	Ile 805	Thr	Tyr	Thr	Thr	Ala 810	Tyr	Leu	Lys	Ala	Ren 815	Tyr
Pro Lys		Trp 820		Ala	Ala	Leu	Leu 825		Cys	Asp	Tyr	Asp 830		Ile
Glu Lys			Lys	Leu	I1e	Gln 840		Ala	His	Ser	Met 845		Ile	Leu
Val Leu 850		Pro	Asp	Ile	Asn 855		Ser	Gly	G1n	Asp 860		Glu	Ala	Thr
Gln Lys 865	Gly	Ile	Arg	Phe 870		Leu	Gly	A1a	Val 875		Gly	Val	Gly	Met 880
Ser Ile	Val	Asp	Ser 885		Val	Glu	G1u	Arg 890	Glu	Lys	Asn	Gly	Pro 895	Tyr
Lys Ser		Gln 900	Asp	Phe	Val	G1n	Arg 905	A1a	Asp	₽he	Lys	Lys 910	Val	Thr
Lys Lys	Gln 915	Leu	Glu	Asn	Leu	Val 920	Asp	Ala	Gly	Thr	Phe 925	Asp	Cys	Phe
Glu Pro 930	Asn	Lys	Asp	Leu	Ala 935	Leu	Ala	Ile	Leu	Asn 940	Asp	Leu	Tyr	Asp
Thr Phe 945				950	_				955					960
Phe Ser	Leu	Asp	Ser 965	Met	A1a	Arg	Asp	Pro 970	Val	Lys	Ile	Thr	Val 975	Ser
Pro Glu		980					985	-				990	-	
	995		_		-	1000)	A1a	His	Pro	Met 1005		Ala	Val
Glu His 1010					1015	5				1020) -	-		
1010 Gly Leu 1025	Pro	His	Gly	Thr 1030	1015 Ile	Ile	Arg	Thr	Val 1035	1020 Phe	Leu	I1e	Asp	Lys 1040
1010 Gly Leu 1025 Val Thr	Pro Thr	His Lys	Gly Ile 1045	Thr 1030 Ser	1015 Ile) Ser	Ile Ala	Arg Glu	Thr Gln 1050	Val 1035 Lys	1020 Phe Lys	Leu Phe	Ile Ala	Asp Leu 1055	Lys 1040 Leu
1010 Gly Leu 1025 Val Thr Gln Val	Pro Thr Ser	His Lys Asp 1060	Gly Ile 1045 Glu	Thr 1030 Ser Val	1015 Ile) Ser Asp	Ile Ala Ser	Arg Glu Tyr 1065	Thr Gln 1050 Glu	Val 1035 Lys) Leu	1020 Phe Lys Pro	Leu Phe Ile	Ile Ala Trp 1070	Asp Leu 1055 Ala	Lys 1040 Leu Asp
Gly Leu 1025 Val Thr Gln Val Met Tyr	Pro Thr Ser Ala 1075	His Lys Asp 1060 Glu	Gly Ile 1045 Glu Tyr	Thr 1030 Ser Val Arg	1015 Ile Ser Asp	Ile Ala Ser Leu 1080	Arg Glu Tyr 1065 Leu	Thr Gln 1050 Glu Glu	Val 1035 Lys) Leu Glu	1020 Phe Lys Pro Asp	Leu Phe Ile Arg	Ile Ala Trp 1070 Leu	Asp Leu 1055 Ala) Ile	Lys 1040 Leu Asp
Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090	Pro Thr Ser Ala 1075 Leu	Hìs Lys Asp 1060 Glu Ala	Gly 11e 1045 Glu Tyr	Thr 1030 Ser Val Arg	1015 Ile Ser Asp Asp Arg 1095	Ile Ala Ser Leu 1080 Arg	Arg Glu Tyr 1065 Leu Ser	Thr Gln 1050 Glu Glu Asp	Val 1035 Lys Leu Glu Ser	1020 Phe Lys Pro Asp Leu 1100	Leu Phe Ile Arg 1085 Arg	Ile Ala Trp 1070 Leu Leu	Asp Leu 1055 Ala) Ile Ser	Lys 1040 Leu Asp Tyr
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp	Pro Thr Ser Ala 1075 Leu Met	His Lys Asp 1060 Glu Ala Arg	Gly Ile 1045 Glu Tyr Ile Asp	Thr 1030 Ser Val Arg Asp Leu 1110	Asp Arg 1095 Ser	Ile Ala Ser Leu 1080 Arg	Arg Glu Tyr 1065 Leu Ser Val	Thr Gln 1050 Glu Glu Asp Asn	Val 1035 Lys Leu Glu Ser Asp 1115	1020 Phe Lys Pro Asp Leu 1100 Ser	Leu Phe Ile Arg 1085 Arg Val	Ile Ala Trp 1070 Leu Leu Leu	Asp Leu 1055 Ala) Ile Ser Ala	Lys 1040 Leu 5 Asp Tyr Cys Glu 1120
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp	Pro Thr Ser Ala 1075 Leu Met	His Lys Asp 1060 Glu Ala Arg Val	Gly Ile 1045 Glu Tyr Ile Asp	Thr 1030 Ser Val Arg Asp Leu 1110 Asp	Asp Arg 1095 Ser Arg Arg Arg	Ile Ala Ser Leu 1080 Arg Thr	Arg Glu Tyr 1065 Leu Ser Val	Glu Glu Glu Asp Asn Ser	Val 1035 Lys Leu Glu Ser Asp 1115 Gln	1020 Phe Lys Pro Asp Leu 1100 Ser	Leu Phe Ile Arg 1085 Arg Val	Ile Ala Trp 1070 Leu Leu Ile Tyr	Asp Leu 1055 Ala Ile Ser Ala Ser 1135	Lys 1040 Leu Asp Tyr Cys Glu 1120 Ser
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp	Pro Thr Ser Ala 1075 Leu Met Glu Lys	His Lys Asp 1060 Glu Ala Arg Val Ser 1140	Gly Ile 1045 Glu Tyr Ile Asp Tyr 1125 Thr	Thr 1030 Ser Val Arg Asp Leu 1110 Asp	1015 Ile Ser Asp Arg 1095 Ser Arg	Ile Ala Ser Leu 1080 Arg Thr Leu Gln	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145	Glu Glu Glu Asp Asn Ser 1130	Val 1035 Lys Leu Glu Ser Asp 1115 Gln Ala	1020 Phe Lys Pro Asp Leu 1100 Ser Lys	Leu Phe Ile Arg 1085 Arg Val Val	Ile Ala Trp 1070 Leu Leu Ile Tyr Lys 1150	Asp Leu 1055 Ala Ile Ser Ala Ser 1135 Lys	Lys 1040 Leu 5 Asp Tyr Cys G1u 1120 Ser 5 Val
Oly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp Thr Lys Glu Thr	Pro Thr Ser A1a 1075 Leu Met G1u Lys Arg 1155	His Lys Asp 1060 Glu Ala Arg Val Ser 1140 Glu	Gly Ile 1045 Glu Tyr Ile Asp Tyr 1125 Thr	Thr 1030 Ser Val Arg Asp Leu 1110 Asp Gly	Asp Arg 1095 Ser Arg Arg Arg Arg	Ile Ala Ser Leu 1080 Arg Thr Leu Gln Val	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145	Gln 1050 Glu Glu Asp Asn Ser 1130 Ser	Val 1035 Lys Leu Glu Ser Asp 1115 Gln Ala	1020 Phe Lys Pro Asp Leu 1100 Ser Lys Met	Deu Phe Ile Arg 1085 Arg Val Val Ile Asp 1165	Ile Ala Trp 1070 Leu Leu Ile Tyr Lys 1150 Leu	Asp Leu 1055 Ala Ile Ser Ala Ser 1135 Lys	Lys 1040 Leu Asp Tyr Cys Glu 1120 Ser Val
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp Thr Lys Glu Thr Leu Arg 1170	Pro Thr Ser A1a 1075 Leu Met G1u Lys Arg 1155 His	His Lys Asp 1060 Glu Ala Arg Val Ser 1140 Glu Ser	Gly Ile 1045 Glu Tyr Ile Asp Tyr 1125 Thr	Thr 1030 Ser Val Arg Asp Leu 1110 Asp Gly Ser Leu	1015 Ile Ser Asp Asp Arg 1095 Ser Arg Ala Pro	Ile Ala Ser Leu 1080 Arg Thr Leu Gln Val 1160	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145 Thr	Glu Asp Asn Ser 1130 Ser Ile	Val 1035 Lys Leu Glu Ser Asp 1115 Gln Ala Ser	1020 Phe Lys Pro Asp Leu 1100 Ser Lys Met Leu Leu 1180	Leu Phe Ile Arg 1085 Arg Val Val Ile Asp 1165	Ile Ala Trp 1070 Leu Leu Ile Tyr Lys 1150 Leu Arg	Asp Leu 105: Ala) Ile Ser Ala Ser 113: Lys) Asn	Lys 1040 Leu Asp Tyr Cys Glu 1120 Ser Val Lys
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp Thr Lys Glu Thr Leu Arg 1170 Ser Gly 1185	Pro Thr Ser Ala 1075 Leu Met Glu Lys Arg 1155 His	His Lys Asp 1060 Glu Ala Arg Val Ser 1140 Glu Ser Glu	Gly Ile 1045 Glu Tyr Ile Asp Tyr 1125 Thr	Thr 1036 Ser Val Arg Asp Leu 1116 Asp Gly Ser Leu Leu 1190	Asp Arg 1095 Ser Arg Arg Arg Arg Arg Arg Arg Arg Arg Ar	Ile Ala Ser Leu 1080 Arg Thr Leu Gln Val 1160 Ile	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145 Thr Leu Val	Glu 1050 Glu Asp Asn Ser 1130 Ser Ile Lys	Val 1035 Lys Leu Glu Ser Asp 1115 Gln Ala Ser Gly Thr	1020 Phe Lys Pro Asp Leu 1100 Ser Lys Met Leu Leu Lys	Deu Phe Ile Arg 1085 Arg Val Ile Asp 1165 Ile Asp 1165 Asp	Ile Ala Trp 107(Leu 5 Leu Ile Tyr Lys 115(Leu Arg Asn	Asp Leu 1055 Ala) Ile Ser Ala Ser 1135 Lys) Asn Lys	Lys 1040 Leu Asp Tyr Cys Glu 1120 Ser Val Lys Tyr
1010 GIY Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp Thr Lys Glu Thr Leu Arg 1170 Ser Gly 1185 Phe Ala	Pro Thr Ser Ala 1075 Leu Met Glu Lys Arg 1155 His	His Lys Asp 1060 Glu Ala Arg Val Ser 1140 Glu Ser Glu	Gly Ile 1045 Glu Tyr Ile Asp Tyr 1125 Thr Ala Ser 1205	Thr 1030 Ser Val Arg Asp Leu 1110 Gly Ser Leu 1190 Pro	Asp Arg 1095 Ser Arg Pro Phe 1175 Ser Asp	Ile Ala Ser Leu 1080 Arg Thr Leu Gln Val 1160 Ile Leu Ala	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145 Thr Leu Val Asp	Thr Gln 1050 Glu Asp Asn Ser 1130 Ser Lys Phe Phe 1210	Val 1035 Leu Glu Ser Asp 1115 Ala Ser Gly Thr 1195 Phe	1020 Phe Lys Pro Asp Leu 1100 Ser Lys Met Leu 1180 Lys Val	Leu Phe Ile Arg 1085 Arg Val Val Ile Asp 1165 Ile Asp Thr	Ile Ala Trp 1070 Leu 5 Leu 5 Leu Tyr Lys 1150 Arg Asn	Asp Leu 105: Ala Ile Ser Ala Ser 113: Lys Asn Lys Gln Asp 121:	Lys 1040 Leu 5 Asp Tyr Cys Glu 1120 Ser 5 Val Lys Tyr Arg 1200 Ile
1010 Gly Leu 1025 Val Thr Gln Val Met Tyr Ala Ile 1090 Arg Trp 1105 Cys Asp Thr Lys Glu Thr Leu Arg 1170 Ser Gly 1185	Pro Thr Ser Ala 1075 Leu Met Glu Lys Arg 1155 His Ser Ser Leu	His Lys Asp 1060 Glu Ala Arg Val Ser 1140 Glu Ser Leu Leu 1220	Gly Ile 1049 Glu Tyr Ile Asp Ile His Ala Ser 1209 Gln	Thr 1030 Ser Val Arg Asp Leu 1110 Gly Ser Leu 1190 Pro	Asp Arg 1095 Ser Arg Pro Phe 1175 Ser Asp	Ile Ala Ser Leu 1080 Arg Thr Leu Gln Val 1160 Ile Leu Ala	Arg Glu Tyr 1065 Leu Ser Val Lys Ser 1145 Thr Leu Val Asp	Thr Gln 1056 Glu Glu Asp Asn Ser 1133 Ser Lys Phe Phe 1210 Thr	Val 1035 Leu Glu Ser Asp 1115 Ala Ser Gly Thr 1195 Phe	1020 Phe Lys Pro Asp Leu 1100 Ser Lys Met Leu 1180 Lys Val	Leu Phe Ile Arg 1085 Arg Val Val Ile Asp 1165 Ile Asp Thr	Ile Ala Trp 1070 Leu 5 Leu 5 Leu Tyr Lys 1150 Arg Asn	Asp Leu 105: Ala Ile Ser Ala Ser 113: Lys Asn Lys Gln Asp 121: Arg	Lys 1040 Leu 5 Asp Tyr Cys Glu 1120 Ser 5 Val Lys Tyr Arg 1200 Ile

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360 Ile Leu Asn Ala Ala Phe Asn Lys Ser Leu Gln Gln Asp Glu Ala Leu 375 380 Arg Ser Gln Leu Glu Lys Arg Ala Tyr Leu Phe Pro Ile Pro Asn Asn 385 390 395 400 Asp Glu Asn Ala Lys Thr Lys Glu Ser Gln Leu Leu Asp Ser Glu Asn 405 410 415 415

Met Val His Tyr Gln His Gln Leu Leu Ser His Leu His Glu Thr Leu 325 330 335 Leu Asp Glu Ala Ile Thr Ala Arg Trp Ser Glu Pro Phe Phe Ile Glu 340 345 350 His Ala Asn Leu Lys Ala Lys Ile Glu Asp Leu Thr Lys Gln Tyr Asp

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Asp Ser Asn Ser Glu Phe Gln Glu Ile Ile Asn Lys Gly Leu Glu Ala 425 Ala Asn Lys Arg Arg Ala Asp Ala Lys Ser Lys Phe Tyr Thr Glu Asp

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Asn Lys Ala Thr Lys Asp Gly Gly Ala Ile Phe Ala Glu Lys Asp Val

295 300 Ser Phe Glu Asn Ile Thr Ser Leu Lys Val Gln Thr Asn Gly Ala Glu 305 310 315 320 Glu Lys Gly Gly Ala Ile Tyr Ala Lys Gly Asp Leu Ser Ile Gln Ser 325 330 335 Ser Lys Gln Ser Leu Phe Asn Ser Asn Tyr Ser Lys Gln Gly Gly Gly 340 345 350 Ala Leu Tyr Val Glu Gly Asp Ile Asn Phe Gln Asp Leu Glu Glu Ile 355 360 365 Arg Ile Lys Tyr Asn Lys Ala Gly Thr Phe Glu Thr Lys Lys Ile Thr 370 375 380 Leu Pro Lys Ala Gln Ala Ser Ala Gly Asn Ala Asp Ala Trp Ala Ser 385 390 400 Ser Ser Pro Gln Ser Gly Ser Gly Ala Thr Thr Val Ser Asn Ser Gly 405 410 415 Asp Ser Ser Ser Gly Ser Asp Ser Asp Thr Ser Glu Thr Val Pro Ala
420
430 Thr Ala Lys Gly Gly Gly Leu Tyr Thr Asp Lys Asn Leu Ser Ile Thr 435 440 445 Asn Ile Thr Gly Ile Ile Glu Ile Ala Asn Asn Lys Ala Thr Asp Val 450 455 460 Gly Gly Gly Ala Tyr Val Lys Gly Thr Leu Thr Cys Glu Asn Ser His 465 475 480 Arg Leu Gln Phe Leu Lys Asn Ser Ser Asp Lys Gln Gly Gly Gle 485 490 495 Tyr Gly Glu Asp Asn Ile Thr Leu Ser Asn Leu Thr Gly Lys Thr Leu
500 505 510 Phe Gln Glu Asn Thr Ala Lys Glu Glu Gly Gly Gly Leu Phe Ile Lys 520 515 525 Ile Asn Asn Thr Ser Glu Lys His Gly Gly Gly Ala Phe Val Thr Lys 545 550 555 560 Glu Ile Ser Gln Thr Tyr Thr Ser Asp Val Glu Thr Ile Pro Gly Ile 565 570 575 Thr Pro Val His Gly Glu Thr Val Ile Thr Gly Asn Lys Ser Thr Gly 580 585 590 Gly Asn Gly Gly Gly Val Cys Thr Lys Arg Leu Ala Leu Ser Asn Leu 595 600 605 Gln Ser Ile Ser Ile Ser Gly Asn Ser Ala Ala Glu Asn Gly Gly Gly 610 615 620 Ala His Thr Cys Pro Asp Ser Phe Pro Thr Ala Asp Thr Ala Glu Gln 625 630 635 Pro Ala Ala Ala Ser Ala Ala Thr Ser Thr Pro Glu Ser Ala Pro Val 645 650 655 Val Ser Thr Ala Leu Ser Thr Pro Ser Ser Ser Thr Val Ser Ser Leu 660 665 670 Thr Leu Leu Ala Ala Ser Ser Gln Ala Ser Pro Ala Thr Ser Asn Lys 675 680 685 Glu Thr Gln Asp Pro Asn Ala Asp Thr Asp Leu Leu Ile Asp Tyr Val Val Asp Thr Thr Ile Ser Lys Asn Thr Ala Lys Lys Gly Gly Gly Ile 705 710 720 Tyr Ala Lys Lys Ala Lys Met Ser Arg Ile Asp Gln Leu Asn Ile Ser 725 730 735 Glu Asn Ser Ala Thr Glu Ile Gly Gly Gly Ile Cys Cys Lys Glu Ser $\frac{740}{740}$ Leu Glu Leu Asp Ala Leu Val Ser Leu Ser Val Thr Glu Asn Leu Val 760 765 755 Gly Lys Glu Gly Gly Gly Leu His Ala Lys Thr Val Asn Ile Ser Asn 770 775 780

Leu Lys Ser Gly Phe Ser Phe Ser Asn Asn Lys Ala Asn Ser Ser Ser 785 790 795 800 Thr Gly Val Ala Thr Thr Ala Ser Ala Pro Ala Ala Ala Ala Ala Ser 810 805 Leu Gln Ala Ala Ala Ala Ala Val Pro Ser Ser Pro Ala Thr Pro Thr 820 825 830 Tyr Ser Gly Val Val Gly Gly Ala Ile Tyr Gly Glu Lys Val Thr Phe 835 840 845 Ser Gln Cys Ser Gly Thr Cys Gln Phe Ser Gly Asn Gln Ala Ile Asp 850 855 860 Asn Asn Fro Ser Gln Ser Ser Leu Asn Val Gln Gly Gly Ala Ile Tyr 865 870 875 880 Ala Lys Thr Ser Leu Ser Ile Gly Ser Ser Asp Ala Gly Thr Ser Tyr 885 890 Ile Phe Ser Gly Asn Ser Val Ser Thr Gly Lys Ser Gln Thr Thr Gly 900 905 910Gln Ile Ala Gly Gly Ala Ile Tyr Ser Pro Thr Val Thr Leu Asn Cys 915 920 · 925 Pro Ala Thr Phe Ser Asn Asn Thr Ala Ser Met Ala Thr Pro Lys Thr 930 935 940 Ser Ser Glu Asp Gly Ser Ser Gly Asn Ser Ile Lys Asp Thr Ile Gly 945 950 955 960 Gly Ala Ile Ala Gly Thr Ala Ile Thr Leu Ser Gly Val Ser Arg Phe 965 970 975 Ser Gly Asn Thr Ala Asp Leu Gly Ala Ala Ile Gly Thr Leu Ala Asn 985 Ala Asn Thr Pro Ser Ala Thr Ser Gly Ser Gln Asn Ser Ile Thr Glu 995 1000 1005 Lys Ile Thr Leu Glu Asn Gly Ser Phe Ile Phe Glu Arg Asn Gln Ala 1010 1015 1020 Asn Lys Arg Gly Ala Ile Tyr Ser Pro Ser Val Ser Ile Lys Gly Asn 1025 1030 1035 1040 Asn Ile Thr Phe Asn Gln Asn Thr Ser Thr His Asp Gly Ser Ala Ile 1045 1050 1055 Tyr Phe Thr Lys Asp Ala Thr Ile Glu Ser Leu Gly Ser Val Leu Phe 1060 1065 1070 Thr Gly Asn Asn Val Thr Ala Thr Gln Ala Ser Ser Ala Thr Ser Gly 1075 1080 1085 Gln Asn Thr Asn Thr Ala Asn Tyr Gly Ala Ala Ile Phe Gly Asp Pro 1090 1095 1100 Gly Thr Thr Gln Ser Ser Gln Thr Asp Ala Ile Leu Thr Leu Leu Ala 1105 1110 1115 1120 Ser Ser Gly Asn Ile Thr Phe Ser Asn Asn Ser Leu Gln Asn Asn Gln 1125 1130 1135 Gly Asp Thr Pro Ala Ser Lys Phe Cys Ser Ile Ala Gly Tyr Val Lys 1140 1145 1150Leu Ser Leu Gln Ala Ala Lys Gly Lys Thr Ile Ser Phe Phe Asp Cys 1155 1160 1165 Val His Thr Ser Thr Lys Lys Ile Gly Ser Thr Gln Asn Val Tyr Glu 1170 1175 1180 Thr Leu Asp Ile Asn Lys Glu Glu Asn Ser Asn Pro Tyr Thr Gly Thr 1185 1190 1195 1200 Ile Val Phe Ser Ser Glu Leu His Glu Asn Lys Ser Tyr Ile Pro Gln 1205 1210 1215 Asn Ala Ile Leu His Asn Gly Thr Leu Val Leu Lys Glu Lys Thr Glu 1220 1225 1230 Leu His Val Val Ser Phe Glu Gln Lys Glu Gly Ser Lys Leu Ile Met 1235 1240 1245 Lys Pro Gly Ala Val Leu Ser Asn Gln Asn Ile Ala Asn Gly Ala Leu 1250 1255 1260 Val Ile Asn Gly Leu Thr Ile Asp Leu Ser Ser Met Gly Thr Pro Gln

Als Gily Gilu Tile Phe Ser Fro Pro Gilu Leu Arg Ille Val Ala Thr Thr 1285 Ser Ser Ala Ser Gily Gily Ser Gily Val Ser Ser Ser Ille Pro Thr Asn 1300 Pro Lys Arg Ille Ser Ala Ala Ala Pro Ser Gily Ser Ala Ala Thr Thr 1310 Pro Lys Arg Ille Ser Ala Ala Ala Pro Ser Gily Ser Ala Ala Thr Thr 1315 1315 Pro Thr Met Ser Gilu Asn Lys Val Phe Leu Thr Gily Asp Leu Thr Leu 1330 1326 Pro Thr Met Ser Gilu Asn Phe Tyr Gin Asn Pro Met Leu Gily Ser Asp 1345 Ille Asp Pro Asn Gily Asn Phe Tyr Gin Asn Pro Met Leu Gily Ser Asp 1335 Leu Asp Val Pro Leu Ille Lys Leu Pro Thr Asn Thr Ser Asp Val 360 Leu Asp Val Pro Leu Ille Lys Leu Pro Thr Asn Thr Ser Asp Val 360 Leu Asp Val Pro Leu Ille Lys Leu Pro Thr Asn Thr Ser Asp Val 360 Leu Asp Val Pro Leu Asp Ser Asn Pro Gin Thr Gily Lys Leu Gin 1370 Met Gily Thr Thr Leu Asp Ser Asn Pro Gin Thr Gily Lys Leu Gin 1395 Met Gily Thr Thr Pro Asp Thr Tyr Arg Arg Trp Val 170 Ille Pro Arg 1470 Asp Asn His Phe Tyr Ala Asn Ser Ille Leu Gily Ser Gin Asn Ser Met 1425 Asp Asn His Phe Tyr Ala Asn Ser Ille Leu Gily Ser Gin Asn Ser Met 1426 Asp Asn His Phe Tyr Asn Asn He Trp Val Ser Gily Val Gily Thr 1460 1445 Phe Asp Asp Ille Ala Tyr Asn Asn Phe Trp Val Ser Gily Val Gily Thr 1475 Net Gily Thr Ser Val Ala 11e Asp Ala Lys Pro Arg Gin Asp Phe 1495 Ser Arg Gily Thr Ser Val Ala 11e Asp Ala Lys Pro Arg Gin Asp Phe 1495 Ille Usily Ala Ala Phe Ser Lys Met Val Gily Lys Thr Lys Ala Ille 1505 Ser Arg Gily Thr Ser Val Ala 11e Asp Ala Lys Pro Arg Gin Asp Phe 1505 Ala Ser Val Tyr Gily Gily Sep Phe Leu Tyr Pe Ser Ille His Gil Arg 155 Ala Ser Val Tyr Gily Gily Sep Phe Leu Tyr Pe Leu Leu Asn Dys Gin 1575 Asn Lys Gily Asp Trp Gilu Asp Leu Gily Thr For Ser Ille His Gilu Arg 1555 Ala Ser Val Tyr Asp Er Pro Arg His Phe Asp Ser Ser Lys Arg Ille Thr 1605 Asn Lys Gily Asp Trp Gilu Asp Leu Gily Try Ser File His Gilu Arg 1505 Asn Lys Gily Asp Trp Gilu Asp Leu Gily Try Ser File His Gilu File Ser Met Asp Leu Lys Gilu Fro Ser Ille Arg Gin Lys Gilu Fro Ser Lie His Gilu Leu Gilu Tyr Ser Pro Arg Gilu Leu Gilu Tyr Se							_				
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Ser Ser Ala Ser Gly Gly Ser Gly Val Ser Ser Ser Ile Pro Thr Asn 1300 1301 1301 1305 1310 1315 1316 1315 1316 1315 1315 1316 1315 1315 1316 1315 1315 1316 1316 1315 1315 1316 1317 1318	ALC GLY GLG			PIO (116	. va.t	ura		
Pro Lys Arg Ile Ser Ala Ala Ala Pro Ser Gly Ser Ala Ala Thr Thr 1315 1320 1325 1326 1330 1335 1340 1335 1340 1335 1340 1355 1356 1355 1356 1355 1356 1355 1356 1355 1356 1355 1356 1355 1356 1360 1365 1370 1365 1370 1365 1370 1365 1370 1375 1380 1385 1370 1375 1380 1385 1390 1385 1390 1385 1390 1385 1390 1380 1385 1390 1405 1395 1395 1395 1360 1405 1395 1395 1400 1405 1395 1400 1405	Ser Ser Ala	Ser Gly			Val S		Ser	Ile		Thr	
Pro Thr Met Ser Glu Asn Lys Val Phe Leu Thr Gly Asp Leu Thr Leu 1330	Pro Lys Arg	Ile Ser	Ala Ala	Ala	Pro S	er Gly	Ser		Ala		Thr
11	Pro Thr Met			Val		eu Thr		Asp		Thr	Leu
Leu Asp Val Pro Leu Ile Lys Leu Pro Thr Asr Thr Ser Asp Val Cln 1365	Ile Asp Pro	Asn Gly	Asn Phe		Gln A		Met		Gly	Ser	
Val Tyr Asp Leu Thr Leu Ser Gly Asp Leu For Gln Lys Gly Tyr 1390 Met Gly Thr Thr Thr Leu Asp Ser Asn Pro Gln Thr Gly Leu Gly Leu Gl 1400 1405 1410 1410 1410 1420 1420 Asp Asp Thr Thr Asp Asp Thr Thr Asp Asp Thr Thr Asp Asp Ser Gln Gly Leu Leu Gly Ser Gln Asp As			Ile Lys	Leu l		hr Asr		Ser	Asp		Gln
Met Gly Thr Trp Thr Leu Asp Ser Asn Pro Gln Thr Gly Lys Leu Gln 1955 1400 1405 1405 1406 1405 1406 1405 1406 1405 1406 1406 1406 1406 1406 1406 1406 1406	Val Tyr Asp	Leu Thr	Leu Ser		Asp I	eu Phe	Pro	Gln		Gly	
Ala Arg Trp Thr Phe Asp Thr Tyr Arg Arg Trp Val Tyr Ile Pro Arg 1410 1410 1415 1410 1415 1415 1425 1440 1445 1455 1440 1445 1455 1440 1445 1455 1450 1445 1450 1455 1460 1445 1460 1445 1460 1445 1460 1445 1460 1465 1460 1467 Phe Leu Ala Gln Glr Jleu Ile Asn Asn Met Leu Asn Asn Ala Arg 1460 1465 1460 1465 1460 1465 1460 1465 1460 1465 1467 Phe Leu Ala Gln Glr Glr Hr Fro Leu Ser Glu Glu Fhe Ser Tyr Tyr 1475 1460 1480 Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Lys Fro Arg Gln Asp Phe 1490 1495 1495 1496 1496 1496 1497 1498	Met Gly Thr	Trp Thr	Leu Asp	Ser A	Asn P	ro Glr	Thr		Lys		Gln
Asp Ran His Phe Tyr Nia Ash Ser Ile Leu Gly Ser Gin Ash Ser Met 1425 14430 1435 14450 Ile Val Val Lys Gin Gly Leu Ile Ash Ash Met Leu Ash Ash Ala Arg 1445 1460 1445 1465 1465 1467 Phe Asp Asp Ile Ala Tyr Ash Ash Phe Trp Val Ser Gly Val Gly Thr 1460 1467 1467 Phe Leu Ala Gin Gly Gly Thr Pro Leu Ser Glu Glu Phe Ser Tyr Tyr 1475 1480 1483 Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Lys Fro Arg Gin Asp Phe 1490 1495 1500 Ile Leu Gly Ala Ala Phe Ser Lys Met Val Gly Lys Thr Lys Ala Ile 1505 1510 1515 1520 Lys Lys Met His Ash Tyr Phe His Lys Gly Ser Glu Tyr Ser Tyr Gin 1505 1550 Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Ash Lys Gln 1550 1550 His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly Gly Els Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1555 His Ile Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1555 Grid Tyr Gly Glu For Ser Lys Asp Ser Ser Ile His Glu Arg 1555 His Ile Lys His Asp Thr Glu Asp Leu Lys Asp Ser Ser Lys Arg Ile Thr 1560 Ser Met Asp Leu Lys Glu Fro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1560 Wal Tyr Gly Glu Glu Tyr Ser Ser Le Arg Gln Lys Gln 1615 Wal Tyr Gly Glu Gly Lys Flee Lav Tyr Leu Ala Sp Leu Arg 1661 Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Ash 1630 Ann Ile Leu Met Tyr Asp Lys Leu Ala Glu Gly Ala Ile Met Asp Ceu 1635 Ann Ile Leu Met Tyr Asp Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile 1665 1660 Ann Ile Leu Met Tyr Asp Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile 1665 1670 Tyr Arg Ash Ash Pro Val Cys Lys Tyr Arg Ala Lu Ser Ser Ash Glu 1707 Sen Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gin Met Thr 1695 Ala Glu Glu The Asp Pro Thr Arg Thr Ser Ala Arg Ala 1700 Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1715 Asp Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gin Met Thr Ser Ala Arg Ala 1700 Clu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1730 Cys Gly Ala Arg Met Ile Phe	Ala Arg Trp		Asp Thr	Tyr i			Val	Tyr		Pro	Arg
11	Asp Asn His	Phe Tyr	Ala Asn		Ile I		Ser		Asn	Ser	
Phe Asp Asp Ile Ala Tyr Asn Asn Phe Trp Val Ser Gly Val Gly Thr 1460 1460 1460 1460 1460 1465 1460 1465 1465 1467 1465 1467 1468 Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Iys Fro Arg Gln Asp Phe 1490 1495 1495 1495 1495 1495 1495 1495 1500 1495 1501 1501 1501 1501 1505 1501 1505 1501 1505 1501 1505 1501 1505 1501 1505 1501 1505 1501 1505 1506 1507 1507 1508 1508 1509 1507 1508 1509			Gly Leu	Ile A		sn Met		Asn	Asn		Arg
Phe Leu Ala Gin Gly fir Pro Leu Ser Glu Glu Phe Ser Tyr Tyr 1475 1480 1485 Ser Arg Gly Thr Ser Val Ala Ile Asp Ala Iys Fro Arg Gin Asp Phe 1490 Ile Leu Gly Ala Ala Phe Ser Lys Met Val Gly Lys Thr Lys Ala Ile 1505 Ile Leu Gly Ala Ala Phe Ser Lys Met Val Gly Lys Thr Lys Ala Ile 1505 1520 Lys Lys Wet His Asn Tyr Fhe His Lys Gly Ser Glu Tyr Ser Tyr Gln 1525 1530 Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Asn Lys Gln 1540 1540 Eis Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly Gly 1550 His Lie Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1575 Asn Lys Gly Asp Trp Glu Bay Leu Lys Gln 1585 Asn Lys Gly Asp Trp Glu Say Leu Lys Asp Ser Ser Lys Arg Ile 1580 Ser Met Asp Leu Lys Glu Fro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1635 Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln 1615 Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Pro Car Lys Asp Ser Ser Lys Arg Inc 1635 Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala In Met Tyr Asp Asn 1635 Asn Ile Leu Met Tyr Asn Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile 1665 Asn Ile Leu Met Tyr Asn Lys Leu Ala Leu Ala Tyr Met Pro Ser Ile 1665 1670 Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1705 Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Fro Thr Arg Thr Ser Ala Arg Ala Glu Glu Fro Fro Thr Arg Thr Ser Ala Arg Ala Glu Tyr Fro Thr Leu Tyr Gly 1715 Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Ser Gin Met Thr Ser Ala Arg Ala Glu Tyr Fro Thr Ile Gly 1725 Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Ser Gin Met Thr Ser Ala Tyr Met Tyr Tyr Arg Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Ser Gin Met Thr Ser Ala Arg Ala Cys Lys Tyr Arg Val Cys Lys Tyr Arg Val Cys Cys Gly Ala Arg Met Ile Tyr Ser Thr Gln Leu Tyr Gly 1725 Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Ser Gin Met Thr Ser Ala Tyr Met Tyr Tyr Lyr Arg Val Arg Ala Arg Met Ile Tyr Ser Thr Gln Leu Tyr Gly 1725 Asn Tyr Thr Ile Asp Val Gly Wet Tyr Thr Leu Ser Gin Met Thr Ser 1730 Cys Gly Ala Arg Met Ile Phe	Phe Asp Asp	Ile Ala			Phe T	rp Val	Ser	Gly		Gly	
Ser Arg GLY Thr Ser Val Ala IIe Asp Ala Lys Pro Arg Gln Asp Phe 1490	Phe Leu Ala	Gln Gln	Gly Thr	Pro 1	Leu S	er Glu	Glu		Ser		Tyr
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Lys Lys Met His Asan Tyr Phe His Lys Gly Ser Glu Tyr Ser Tyr Gln 1525 1530 Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Phe Leu Leu Asan Lys Gln 1540 His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly 1555 His Is Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1570 Asan Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg 1180 Asan Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 Asan Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 Asan Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 Asan Lys Gly Leu Glu Tro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1630 Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Asan 1620 Glu Ile Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Asan 1635 Leu Ser Leu Fro Val Gly Cys Ala Val Glu Gly Ala Ile Ber Leu 1670 Tyr Arg Asan Asan Pro Val Cys Leu Ala Leu Ala 1670 Tyr Arg Asan Asan Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asan Glu 1685 1690 Ala Glu Gln Val Ile Cys Gly Val Pro Thr Arg Thr Ser Ala Arg Ala 1700 Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1725 Asan Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr Ser 1730 Cys Gly Ala Arg Met His Phe	Ile Leu Gly	Ala Ala	Phe Ser		Met V		Lys		Lys	Ala	
Ala Ser Val Tyr Gly Gly Lys Phe Leu Tyr Fhe Leu Leu Asn Lys Gln 1540 His Gly Trp Ala Leu Pro Phe Leu Ile Gln Gly Val Val Ser Tyr Gly 1555 His Ile Lys His Asp Thr Thr Thr Leu Tyr Pro Ser Ile His Glu Arg 1570 Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg 1580 Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg 11850 Ser Met Asp Leu Lys Glu Pro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1610 Val Tyr Gly Glu Leu Glu Tyr Ser Ser Ile Arg Gln Lys Gln Pro Thr Thr Thr Leu Tyr Gly Glu Fro Tyr Asp Fro Arg His Pro Arg Asn 1620 Glu Lle Asp Tyr Asp Pro Arg His Phe Asp Asp Cys Ala Tyr Arg Asn 1657 Leu Ser Leu Fro Val Gly CysAla Val Glu Gly Ala Ile Met Asn Cys 1650 And The Leu Met Tyr Asn 192 Leu Ala Arg Asn 1680 Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1685 Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1690 Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1725 Asn Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr Ser Ala Arg Ala Tyn Tyr Arg Val Ala Arg Met Ile Phe		His Asn	Tyr Phe	His 1	Lys G	ly Ser		Tyr	Ser	Tyr 1535	Gln
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His Ile Lys His Asp Thr Thr Thr Lou Tyr Pro Ser Ile His Glu Arg 1570 Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585 Asn Lys Gly Asp Trp Glu Asp Leu Gly Trp Leu Ala Asp Leu Arg Ile 1585: 1660 Ser Met Asp Leu Lys Glu Pro Ser Lys Asp Ser Ser Lys Arg Ile Thr 1615 Val Tyr Gly Glu Euo Glu Tyr Ser Ser Ile Arg Gln Lys Gln Phe Thr 1620 Glu Lleu Sel Leu Ser Ser Asp His She Asp Asp Cys Ala Tyr Arg Asn 1625 Glu Lle Asp Tyr Asp Pro Arg His She Asp Asp Cys Ala Tyr Arg Asn 1615 Leu Ser Leu Pro Val Gly Cys Ala Val Glu Gly Ala Tie Met Asn Cys 1635 Ala 1616 Leu Ser Leu Met Tyr Asn Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1645 Tyr Arg Asn Asn Pro Val Cys Lys Tyr Arg Val Leu Ser Ser Asn Glu 1685 Ala Gly Gln Val Ile Cys Gly Val Pro Thr Arg Thr Ser Ala Asq Ala 1706 Glu Tyr Ser Thr Gln Leu Tyr Leu Gly Pro Phe Trp Thr Leu Tyr Gly 1725 Asn Tyr Thr Ile Asp Val Gly Met Tyr Thr Leu Ser Gln Met Thr Ser 1730 Cys Gly Ala Arg Met Ile Phe			Pro Phe	Leu :	Ile G	ln Gly	Val		Ser		Gly
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Met	Ser	Trp	Val 340	Lys	Asn	Ile	Val	Asp 345	Pro	Ser	Glu	Val	Val 350	Asn	Lys
Gly	Asp	Glu 355	Val	Glu	Ala	Ile	Val 360	Leu	Ser	Ile	Gln	Lys 365	Asp	Glu	Gly
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Ile 385	Glu	Glu	Lys	Tyr	Pro 390	Ile	Gly	Leu	His	Val 395	Asn	Ala	Glu	Ile	Lys 400
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Pro Asp Glu Lys Cys Ser Val Tyr Pro Leu Glu Asp Gly Thr Val Lys 305 310 315 320

Val Ile Phe Asp Val Pro Val Lys Ala Val Thr Pro Gly Gln Thr Val 325 330 335

Pro Met Ile His Gln Leu 355